# **GOVERNMENT ARTS COLLEGE (AUTONOMOUS)**

(Affiliated to Bharathidasan University, Tiruchirappalli – 620 024) **KUMBAKONAM – 612 002 TAMILNADU** 

# Syllabusfor M.Sc. BIOCHEMISTRY

Choice Based Credit System (CBCS) (With effect from 2023 - 2024 onwards)



# Department of Biochemistry Government Arts College (Autonomous)

(Affiliated to Bharathidasan University, Tiruchirappalli – 620 024)

Kumbakonam – 612 002 Tamilnadu

# GOVERNMENT ARTS COLLEGE (AUTONOMOUS), KUMBAKONAM – 612 002 Course Structure under CBCS for Science (2023 -2024 Onwards)

		SYLLABUS FOR M.Sc BIOCHEMISTI			ARDS)			
~		SCHEME OF STUDY AND			1			ı
Semes- ter	Course Code	Title of the Paper	Credits	Inst.Hrs/ Week	Exam Hours	Ma Int.	rks Ext.	Total
I	23P1B1	Core Course – I: Basics of Biochemistry	5	6	3	25	75	100
-	23P1B2	Core Course – II: Biochemical and	5	6	3	25	75	100
		Molecular Biology Techniques						
	23P1BP1	4	6	6	40	60	100	
		Core Practical – I: Laboratory Course on Biomolecules and Biochemical						
		Techniques						
	23P1B3EC	Elective Course– I: Physiology and Cell Biology (To include Hormones)	3	6	3	25	75	100
	23P1B4EC	Elective Course– II: Microbiology and	3	6	3	25	75	100
	23F1B4EC	Immunology	3	0	3	23	13	100
		Total	20	30	_	Total I	l Marks	500
II	23P2B5	Core Course – III: Enzymology	5	6	3	25	75	100
	23P2B6	Core Course – IV: Cellular Metabolism	5	6	3	25	75	100
	23P2BP2	Core Practical – II: Laboratory Course	4	6	6	40	60	100
	23F2DF2	on Enzymology, Microbiology and Cell Biology	4	0	0	40	00	100
	23P2B7EC	Elective Course– III: Molecular Biology	3	4	3	25	75	100
	23P2B8EC	Elective Course– IV: Energy and Drug	3	4	3	25	75	100
		Metabolism						
	23P2B9SEC	Skill Enhancement Course [SEC] – I: Nutritional Biochemistry	2	4	3	25	75	100
		Total	22	30	_	Total M	[arks	600
III	23P3B10	Core Course – V: Clinical Biochemistry	5	6	3	25	75	100
	23P3B11	Core Course – VI: Gene Editing, Cell	5	6	3	25	75	100
		and Gene Therapy						
	23P3B12	Core Course – VII / Industrial Module:	5	6	3	25	75	100
		Industrial Microbiology			_			
	23P3BP3	Core Practical – III: Laboratory Course	4	6	6	40	60	100
	23P3B13EC	on Clinical Biochemistry  Elective Course– V: Biostatistics and	4	3	3	25	75	100
	23F 3B13EC	Data Science	4	3	3	23	13	100
	23P3B14SEC	Skill Enhancement Course [SEC] – II:	2	3	3	25	75	100
		Molecular Basis of Diseases and						
		Therapeutic Strategies						
		Internship / Industrial Activity	2	-	-	-	-	-
W. 2204015 C C		Total	27	30	-	Total I		600
IV	23P4B15	Core Course – VIII: Pharmaceutical Biochemistry	5	6	3	25	75	100
	23P4BP4	Core Practical – IV: Laboratory Course	5	6	6	40	60	100
		on Nutritional Biochemistry and Biology						
	23P4B16EC	Elective – VI: Biochemical Toxicology	4	4	3	25	75	100
	23P4B17SEC	Skill Enhancement Course [SEC] – III:	2	4	3	25	75	100
		Biosafety, Lab Safety and IPR						
	23P4B18PW	Project Work with Viva - Voce	7	10	-	20	80	100
	23P4BEA	Extension Activities	1	-	-	-	-	-
		Total	24	30	-	Total		500
		Net Total	93	120	-	Net 7		2200
					]	Ma	rks	

# **M.Sc - Biochemistry Course Structure**

Course	Nos.	Credits	Marks
Core Course	8	40	800
Core Practical	4	17	400
Elective Course	6	18	600
Skill Enhancement Course [SEC]	3	6	300
Project Work with Viva - Voce	1	7	100
Internship / Industrial Activity		2	
Extension Activity		1	
Total	22	91	2200

# **Question Paper Pattern for Semester Theory Examination**

Sl. No.	Sl. No. Section Marks Number of Questions Marks					
1	Section A	2 marks	Two questions from each unit	$10 \times 2 = 20$	20	
2	2   Section B   5 marks   One SET from each unit –   5 x 5				25	
			Either Or type			
3	Section C	10 marks	One question from each unit –	One question from each unit $-$ 3 x 10 = 30		
	Answer any THREEquestions					
	75					
	25					
	100					

# **Question Paper Pattern**

# Question paper pattern for semester practical examination

Internal Maximum: 40 External Maximum: 60

# **Project Evaluation**

Maximum Marks: 100

Project Work with	Report Maximum: 80	Viva Voce Maximum: 20
Viva - Voce	To be evaluated for overall objective and	Performance of the candidate
viva - voce	quality of work presented in the report.	remormance of the candidate

Programme:	M.Sc - BIOCHEMISTRY
Programme Code:	LIFC
<b>Duration:</b>	2 years
Programme Outcomes:	<ul> <li>PO1: To make students understand the importance of biochemistry as a subject that deals with life processes, as well as the concepts, theories and experimental approaches followed in biochemistry, in order to pursue a research career, either in an industry or academic setting.</li> <li>PO2: To develop analytical and problem-solving skills.</li> <li>PO3: To create awareness among the students on the interconnection between the interdisciplinary areas of biochemistry.</li> <li>PO4: To give the necessary practical skills required for biochemical techniques and analysis.</li> <li>PO5: To develop a communication and writing skills in students.</li> <li>PO6: To develop leadership and teamwork skills.</li> <li>PO7: To emphasize the importance of good academic and work ethics and their social implications.</li> <li>PO8: To emphasize the importance of continuous learning and to promote lifelong learning and career development.</li> <li>PO9: To teach students how to retrieve information from a variety of sources, including libraries, databases and the internet.</li> <li>PO10:To teach students to identify, design and execute a research problem,</li> </ul>
	analyze and interpret data and learn time and resource management.
Programme	On successful completion of this course, students should be able to:
Specific Outcomes (PSO):	PSO1: Understand the principles and methods of various techniques in Biochemistry, Immunology, Microbiology, Enzyme kinetics and Molecular Cell Biology. Based on their understanding, the students may would be able to design and execute experiments during their final semester project, and further research programs.
	PSO2: Insight on the structure-function relationship of biomolecules, their synthesis and breakdown, the regulation of these pathways, and their importance in terms of clinical correlation. Students will also acquire knowledge of the principles of nutritional biochemistry and also understand diseases and their prevention.  PSO3: To understand the concepts of cellular signal transduction pathways and the association of aberrant signal processes with various diseases. Acquire insight into the immune system and its responses, and use this knowledge in the processes of immunization, vaccine development, transplantation and organ rejection.
	<ul> <li>PSO4: To visualize and appreciate the central dogma of molecular biology, regulation of gene expression, molecular techniques used in rDNA technology, gene knocks-out and knock-in techniques.</li> <li>PSO5: To create awareness in students about the importance of good laboratory practices and the importance of ethical and social responsibilities of a researcher. Teach them how to review literature and the art of designing and executing experiments independently and also work as a part of a team.</li> </ul>

<b>Course / Course Code:</b>	CORECOURSE –I / 23P1B1
TitleoftheCourse:	BASICS OF BIOCHEMISTRY
Semester / Credits:	I/5
Pre-requisites, ifany:	BasicKnowledgeofBiochemistry and Biomolecules
CourseObjectives:	Themainobjectivesofthiscourseareto:
	1. Students will be introduced to the structure of biomolecules.
	2. The significance of carbohydrates in biological processes will be
	understood.
	3. The structure, properties and biological significance of lipids in the
	biological system will be studied
	4. Students will learn about the concepts of protein structure and their
	significance in biological processes and creatively
	comprehendtheroleof membranecomponents with
	theirbiologicalsignificance.
	5. Students will gain knowledge about
	thestructuresandfunctionalrolesofnucleicacids in the
	biologicalsystem.
CourseOutcomes:	On successful completionofthecourse, the students should be able to:
	CO1: Explain the
	chemicalstructureandfunctionsofcarbohydrates(K1,K2).
	CO2: Usingtheknowledgeoflipidstructureandfunction, explain how it
	plays aroleinSignalingpathways (K3,K4).
	CO3: Describe the various levels of structural organisation of proteins
	andtheroleof proteinsinbiologicalsystem(K4,K5).
	CO4: Applytheknowledgeofproteinsincell-cellinteractions(K3,K4).
	CO5.
	Applyingtheknowledgeofnucleicacidsequencinginresearchanddiag
	nosis(K2,K3&K4).

	Units
_	Carbohydrates- Classification, structure (configurations and conformations, anomeric
1	forms), function and properties of monosaccharides, mutarotation, Disaccharides and oligosaccharides with suitable examples. Polysaccharides - Homopolysaccharides
	(starch, glycogen, cellulose, inulin, dextrin, agar, pectin, dextran). Heteropolysaccharides
	- Glycosaminoglycans- source, structure, functions of hyaluronic acid, chondroitin
	sulphates, heparin, keratan sulphate. Glycoproteins - proteoglycans. O- Linked and N-linked glycoproteins. Biological significance of glycan. Blood group polysaccharides.
	Bacterial cell wall (peptidoglycans, teichoic acid) and plant cell wall carbohydrates.

- II Lipids - Classification of lipids, structure, properties and functions of fatty acids, triacylglycerols, phospholipids, glycolipids, sphingolipids and steroids - Biological importance. Eicosanoids- classification, structure and functions of prostaglandins, thromboxanes, leukotrienes. Lipoproteins – Classification, structure, transport (endogenous and exogenous pathway) and their biological significance. Ш Overview of Aminoacids - classification, structure and properties of amino acids, Biological role. Non Protein aminoacids and their biological significance. Proteins classification based on composition, structure and functions. Primary, secondary, super secondary (motifs) (Helix-turn –helix, helix-loop-helix, Beta-alpha-beta motif, Rosemann Rossmann fold, Greek key),tertiary and quaternary structure of proteins. Structural characteristics of collagen and hemoglobin. Determination of sequence. Chemical synthesis of a peptide, Forces involved in stabilization of protein structure. Ramachandran plot. Folding of proteins. Molecular chaperons – Hsp 70 and Hsp 90 - biological role. IVMembrane Proteins - Types and their significance. Cytoskeleton proteins - actin, tubulin, intermediate filaments. Biological role of cytoskeletal proteins. Membrane structure-fluid mosaic model.  $\mathbf{V}$ Nucleic acids – types and forms (A, B, C and Z) of DNA. Watson-Crick model-Primary, secondary and tertiary structures of DNA. Triple helix and quadruplex DNA. Mitochondrial and chloroplast DNA. DNA super coiling (calculation of Writhe, linking and twist number). Determination of nucleic acid sequences by Maxam Gilbert and Sanger's methods. Forces stabilizing nucleic acid structure. Properties of DNA and RNA. C-value, C-value paradox, Cot curve. Structure and role of nucleotides in cellular communications. Major and minor classes of RNA, their structure and biological functions. Reading List (PrintandOnline)
  - 1. https://bio.libretexts.org/Bookshelves/Biochemistry/Book%3A\_Biochemistry\_Online\_(Jakubowski)
  - 2. https://www.thermofisher.com/in/en/home/life-science/protein-biology/protein-biology-learning-center/protein-biology-resource- library/pierce-protein-methods/protein-glycosylation.html
  - 3. https://ocw.mit.edu/courses/biology/7-88j-protein-folding-and- human-disease-spring-2015/study-materials/
  - 4. https://www.open.edu/openlearn/science-maths- technology/science/biology/nucleic-acids-and-chromatin/content-section-3.4.2
  - 5. https://www.genome.gov/genetics-glossary/Cell-Membrane
  - 6.https://nptel.ac.in/content/storage2/courses/102103012/pdf/mod3.pdf

### SelfStudy

- 1. Classification of sugars
- 2. Nutritional classification of fatty acids

### RecommendedTexts

- 1. DavidL.NelsonandMichaelM.Cox(2012)LehningerPrinciplesofBiochemistry(6thed)W. H Freeman
- 2. Voet.D&Voet.J.G(2010)Biochemistry(4thed),JohnWiley&Sons,Inc.
- 3. Metzler D.E(2003). The chemical reactions of living cells (2nded), Academic Press.
- 4. ZubayG.L(1999)Biochemistry(4thed),McGrew-Hill.
- 5. Lubert Stryer(2010)Biochemistry(7thed), W.H. Freeman
- 6. Satyanarayan,U(2014)Biochemistry(4thed), ArunabhaSenBooks &Allied(P)Ltd,Kolkata.

#### Method of Evaluation:

Test I	Test II	Assignment	Seminar	Seminar End Semester Examination	
5	10	5	5	75	100

### **Methods of Assessment:**

**Recall(K1)-**Simpledefinitions,MCQ,Recallsteps,Conceptdefinitions.

 $\label{lem:comprehend} \textbf{Understand/Comprehend(K2)-} MCQ, True/False, Shortessays, Concept explanations, shortsummary or overview.$ 

**Application**(**K3**)-Suggestidea/conceptwithexamples,Solveproblems,Observe,Explain.

**Analyse**(**K4**) – Problem-saving questions, Finish a procedure in many steps, Differentiate between various ideas.

**Evaluate**(**K5**)-Longer essay/ Evaluationessay, Critique or justify with prosand cons.

Create (K6) – Check knowledge in specific or offbeat situations, Discussion.

### **Mapping with Programme Outcomes:**

	PO 1	PO 2	PO 3	PO 4	PO 5	PO 6	PO 7	PO 8	PO 9	PO 10
CO1	S	L	M	S	M	M	M	S	M	M
CO2	S	M	L	S	M	M	M	S	M	M
CO3	S	M	M	S	S	M	L	S	M	M
CO4	S	M	M	S	M	M	M	S	M	M
CO5	S	S	M	S	S	M	M	S	M	M

S- Strong M-Medium L-Low

Course / Course Code:	CORECOURSE -II / 23P1B2					
Title of the Course:	BIOCHEMICAL AND MOLECULAR BIOLOGY TECHNIQUES					
Semester / Credits:	I/5					
Pre-requisites, if any:	Comprehensive Knowledge of Tools of Biochemistry/Molecular Biology					
Course Objectives:	Biochemical techniques combine various inter-disciplinary methods in biological research and the course aims to provide students with the following objectives:  1. To understand the various techniques used in biochemical investigation and microscopy.  2. To explain chromatographic techniques and their applications  3. To explain electrophoretic techniques.  4. To comprehend the spectroscopic techniques and demonstrate their applications in biochemical investigations.					
Course Outcomes:	<ul> <li>5. To acquire knowledge of radio labeling techniques and centrifugation.</li> <li>On successful completionofthecourse, students should be able to:</li> <li>CO1: Attain good knowledge in modern used in biochemical investigation and microscopy and apply the experimental protocols to plan and carry out simple investigations in biological research(K1, K5).</li> <li>CO2:Demonstrate knowledge to implement the theoretical basis of chromatography in upcoming practical course work(K3, K5).</li> <li>CO3:Demonstrate knowledge to implement the theoretical basis of electrophoretic techniques in research work(K3, K5).</li> <li>CO4:Tackle more advanced and specialized spectroscopic techniques that are pertinent to research(K1, K2 &amp; K5).</li> <li>CO5:Tackle more advanced and specialized radioisotope and centrifugation techniques that are pertinent to research work(K1, K2 &amp; K5).</li> </ul>					
	Units					
techniques. Orga techniques, cell Biosensors- prin microscope, dark Principle, instrum	General approaches to biochemical investigation, cell culture techniques and microscopic techniques. Organ and tissue slice technique, cell distribution and homogenization techniques, cell sorting, and cell counting, tissue Culture techniques. Cryopreservation, Biosensors- principle and applications. Principle, working and applications of light microscope, dark field, phase contrast and fluorescent microscope. Electron microscope-Principle, instrumentation of TEM and SEM, Specimen preparation and applications-shadow					
Basic principles Chromatography Hydroxy apatite chromatography. development, dete instrumentation, pressure liquid ch	casting, negative staining and freeze fracturing.  Chromatographic Techniques:  Basic principles of chromatography- adsorption and partition techniques. Chiral Chromatography and counter current Chromatography. Adsorption Chromatography – Hydroxy apatite chromatography and hydrophobic interaction Chromatography. Affinity chromatography. Gas liquid chromatography- principle, instrumentation, column development, detectors and applications. Low pressure column chromatography – principle, instrumentation, column packing, detection, quantitation and column efficiency, High pressure liquid chromatography- principle, instrumentation, delivery pump, sample injection unit, column packing, development, detection and application. Reverse HPLC, capillary					

electro chromatography and perfusion chromatography.

### **III** Electrophoretic Techniques:

General principles of electrophoresis, supporting medium, factors affecting electrophoresis, Isoelectric focusing-principle, ampholyte, development of pH gradient and application. PAGE-gel casting-horizontal, vertical, slab gels, sample application, detection-staining using CBB, silver, fluorescent stains. SDS PAGE-principle and application in molecular weight determination principle of disc gel electrophoresis,2D PAGE. Electrophoresis of nucleic acids-agarose gel electrophoresis of DNA, pulsed field gel electrophoresis- principle, apparatus, applications. Electrophoresis of RNA, curve. Microchip electrophoresis and 2D electrophoresis, Capillary electrophoresis.

### **IV** | Spectroscopic Techniques:

Basic laws of light absorption- principle, instrumentation and applications of UV-Visible, IR, ESR, NMR, Mass spectroscopy, Turbidimetry and Nephelometry. Luminometry (Luciferase system, chemiluminescence). X - ray diffraction. Atomic absorption spectroscopy - principle and applications - Determination of trace elements.

### V Radiolabeling Techniques and Centrifugation:

Nature of radioactivity-detection and measurement of radioactivity, methods based upon ionisation (GM counter) and excitation (scintillation counter), autoradiography and applications of radioactive isotopes, Biological hazards of radiation and safety measures in handling radioactive isotopes.

Basic principles of Centrifugation. Preparative ultracentrifugation - Differential centrifugation, Density gradient centrifugation. Analytical ultracentrifugation - Molecular weight determination.

### **Reading List (Print and Online)**

Principles and techniques of biochemistry and molecular biology:

1. https://www.kau.edu.sa/Files/0017514/Subjects/principals%20and%20techiniques%20of%20biochemistry%20and%20molecular%20biology%207th%20ed%

### **Self Study**

- 1. Types of rotors
- 2. Colorimetry principle and applications

### **Recommended Texts**

- 1. Keith Wilson, John Walker (2010) Principles and Techniques of Biochemistry and Molecular Biology (7th ed) Cambridge University Press.
- 2.David Sheehan (2009), Physical Biochemistry: Principles and Applications (2nd ed), Wiley-Blackwell.
- 3.David M. Freifelder (1982) Physical Biochemistry: Applications to Biochemistry and Molecular Biology, W.H. Freeman.
- 4.Rodney F.Boyer (2012), Biochemistry Laboratory: Modern Theory and techniques(2nd ed), Prentice Hall.
- 5. Kaloch Rajan (2011), Analytical techniques in Biochemistry and Molecular Biology, Springer.
- 6. Segel I.H (1976) Biochemical Calculations (2nd ed), John Wiley and Sons.
- 7. Robyt JF (2015) Biochemical techniques: Theory and Practice (1st ed), CBS Publishers & Distributors.

### **Method of Evaluation:**

Test I	Test II	Assignment	Seminar	End Semester Examination	Total	
5	10	5	5	75	100	

### **Methods of Assessment:**

Recall (K1) - Simple definitions, MCQ, Recall steps, Concept definitions.

**Understand/ Comprehend (K2) -** MCQ, True/False, Short essays, Concept explanations, short summary or overview.

Application (K3) - Suggest idea/concept with examples, Solve problems, Observe, Explain.

**Analyse**(**K4**) – Problem-saving questions, Finish a procedure in many steps, Differentiate between various ideas.

Evaluate (K5) - Longer essay/ Evaluation essay, Critique or justify with pros and cons.

Create (K6) – Check knowledge in specific or offbeat situations, Discussion.

### **Mapping with Programme Outcomes:**

	PO 1	PO 2	PO 3	PO 4	PO 5	PO 6	PO 7	PO 8	PO 9	PO 10
CO 1	S	L	M	S	S	L	L	S	S	M
CO 2	S	M	M	S	M	L	M	S	S	L
CO 3	S	M	L	S	M	M	M	S	M	L
CO 4	S	S	L	S	S	M	M	S	M	M
CO 5	S	S	M	S	M	M	M	S	M	M

S - Strong M - Medium L - Low

Course / Course Code:	COREPRACTICAL - I / 23P1BP1
TitleoftheCourse:	LABORATORY COURSE ON BIOMOLECULES AND BIOCHEMICAL TECHNIQUES
Semester / Credits:	I/4
Pre-requisites, If any:	Knowledge on basic principles, Instrumentation of Biochemical
	techniques and metabolic reactions.
Course Objectives:	<ol> <li>To instill skill in students enabling them to apprehend the wider knowledge about principles and techniques to be employed for the biomolecules under investigation.</li> <li>To inculcate the knowledge of various isolation and purification techniques of macromolecules like DNA, RNA, Glycogen and Starch,</li> <li>To perform colorimetric estimations to quantify important metabolites like lactate and tryptophan and minerals like calcium and iron from various sources.</li> <li>To achieve training in subcellular fractionation and to identify them by markers.</li> <li>To achieve training in various chromatographic techniques.</li> <li>To perform the isolation and identification of the organelles of a cell using differential centrifugation.</li> </ol>
	7. To perform phytochemical screening and quantification enabling them to give an insight on phytochemicals this will be useful for future research.
Course Outcomes:	<ul> <li>On successful completionofthecourse, students should be able to: CO1: The student will be able to acquire knowledge and skill in the techniques used in the isolation, purification and estimation of different biomolecules that are widely employed in research (K1, K2&amp; K4).</li> <li>CO2: The students will get acquainted with Principle, Instrumentation and method of Performing UV absorption studies of DNA, Protein and interpreting the alteration occurred during the process of denaturation (K1, K2, K3&amp; K4).</li> <li>CO3: The student will be fine-tune in handling the instruments like colorimeter, spectrophotometer and will be able to estimate the biomolecules and minerals from the given samples (K1, K2&amp; K4).</li> <li>CO4: The student, in addition to acquiring skill in performing various biochemical techniques can also learn to detect presence of phytochemicals and quantify them in the plant sample (K1, K2, K3, K4 &amp; K6).</li> <li>CO5: The students will develop skill in analytical techniques like subcellular fractionation, Paper, Column and Thin layer Chromatography and the group experiments will enable them to build learning skills like team work, Problem solving, Communication ability (K1, K2, K3, K4 &amp; K6).</li> <li>Units</li> </ul>

I	Biochemical studies and estimation of macromolecules						
1	Estimation of Ascorbic Acid						
	Isolation and estimation of DNA from animal tissue.						
	Isolation and estimation of RNA from yeast.						
	Purification of Polysaccharides –Starch and assessment of its purity						
II	Acid Number of oil						
	Iodine number of oil						
	Saponification of oil						
	Estimation of Amino Acid by Formal Titration						
III	Colorimetric estimations						
	Estimation of Protein						
	Estimation of Sugar						
IV	Estimation of minerals						
	Estimation of calcium						
	Estimation of iron						
V	Plant Biochemistry						
	Qualitative analysis - Phytochemical screening						
VI	Group Experiments						
	Separation of identification of lipids by thin layer chromatography						
	Separation of plant pigments from leaves by column chromatography						
	Identification of Sugars by Paper Chromatography						
	Identification of Amino acids by Paper Chromatography						
	Estimation of Flavonoids - Quantitative analysis						
	Reading List (Print and Online)						
	1.https://www.researchgate.net/publication/313745155_Practical_Bio						
	chemistry_A_Student_Companion						
	2.https://doi.org/10.1186/s13020-018-0177-x						
	3.https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5368116/						
	4.https://www.life.illinois.edu/biochem/455/Lab%20exercises/2Photometry/spectropho						
	tometry.pdf						
	5.https://ijpsr.com/bft-article/determination-of-total-flavonoid-and-phenol-content-in-						
	mimusops-elengi-linn/?view=fulltext						
	6.https://skyfox.co/wp-content/uploads/2020/12/Practical-Manual-of-Biochemistry.pdf						
	SelfStudy						
	1. Laboratory Safety Rules, Requirements and Regulations.						
	2. Preparation of standard solutions and reagent						
	3. Denaturation of DNA and absorption studies at 260nm.						
	4. Denaturation of Protein and absorption studies at 280nm  5. Fractionation of sub-callular organization with differential contributation						
	5. Fractionation of sub-cellular organelles by differential centrifugation -						
	Mitochondria and nucleus  6. Identification of the congreted sub-callular fractions using marker engages (env. eng.)						
	6. Identification of the separated sub-cellular fractions using marker enzymes (any one)						
	Books Recommended						
	1. David Plummer (2001) An Introduction to Practical Biochemistry (3rd ed) McGraw						
	Hill Education (India) Private Ltd						

- 2. Jayaraman, J (2011) Laboratory Manual in Biochemistry, New age publishers
- 3. Varley H (2006) Practical Clinical Biochemistry (6th ed), CBS Publishers
- 4. O. Debiyi and F. A. Sofowora, (1978) "Phytochemical screening of medical plants" Iloyidia, vol. 3, pp. 234–246,
- 5. Prof. Sarin A. Chavhan, Prof. Sushilkumar A. Shinde (2019) A Guide to Chromatography Techniques Edition:1
- 6. Analytical techniques in Biochemistry and Molecular Biology; Katoch, Rajan. Springer (2011)

### **Method of Evaluation:**

Test I	Test II	End Semester Examination	Total	Grade
20	20	60	100	

### **Methods of Assessment:**

**Recall (K1) - Simple definitions, MCQ, Recall steps, Concept definitions.** 

**Understand/ Comprehend (K2) - MCQ**, True/False, Short essays, Concept explanations, Short summary or overview.

**Application** (**K3**) - Suggest idea/concept with examples, Solve problems, Observe, Explain.

**Analyse** (**K4**) - Problem-solving questions, Finish a procedure in many steps, Differentiate between various ideas.

Evaluate (K5) - Longer essay/ Evaluation essay, Critique or justify with pros and cons.

Create (K6) - Check knowledge in specific or offbeat situations, Discussion.

### **Mapping with Programme Outcomes:**

	PO 1	PO 2	PO 3	PO 4	PO 5	PO 6	PO 7	PO 8	PO 9	PO 10
<b>CO 1</b>	S	S	S	S	M	S	L	S	M	S
CO 2	S	S	S	S	M	S	L	S	M	S
CO 3	S	S	S	S	M	S	M	S	M	S
CO 4	S	S	S	S	S	S	S	S	S	S
CO 5	S	S	S	S	S	S	S	S	S	S

S - Strong M - Medium L - Low

Course	/ Course Code:	ELECTIVECOURSE - I/ 23P1B3EC					
Title	of the Course:	PHYSIOLOGY AND CELL BIOLOGY					
Seme	ster / Credits:	1/3					
Pre-re	quisites, if any:	Anatomy, Cells and Biological Compounds					
Cour	sa Objectives:	To understand the functions and activities of organs, tissues or cells					
Course Objectives:		and of physical and chemical phenomena involved in the human body.					
Cour	se Outcomes:	On successful completionofthecourse, students should be able to: CO1:Specifically understand the biological and chemical processes within a human cell (K1, K2, K5& K6).					
		CO2:Identify and prevent diseases(K2, K3,K4, k5& K6).					
		CO3:Understand defects in digestion, nutritional deficiencies and					
		intolerances, and gastrointestinal pathologies(K1, K2, K3,K4, K5& K6).					
		CO4:Identify general characteristics in individuals with imbalances of acid- base, fluid and electrolytes(K1, K2, K3,K4, K5& K6).					
		CO5:Process the mechanism: the transmission of biochemical information between cell membrane and nucleus(K1, K2&K5).					
		Units					
I	Major classes of	of cell junctions- anchoring, tight and gap junctions. Major families of					
	cell adhesion n	nolecules (CAMs)- cadherins, integrins. Types of tissues. Epithelium-					
	organisation ar	nd types. The basement membrane. Cell cycle- mitosis and meiosis,					
		ses and regulation. Cell death mechanisms- an overview-apoptosis,					
	necrosis.						
II	_	em- structure and functions of different components of digestive					
		on and absorption of carbohydrates, lipids and proteins, role of bile					
		on and absorption, mechanism of HCl formation in stomach, role of					
	=	es and hormones involved in digestive system. Composition of blood,					
	lymph and CSF. Blood cells - WBC, RBC and energy metabolism of RBC, Blood clotting mechanism and blood groups- ABO and Rhesus system.						
III		stem-Gaseous transport and acid-base homeostasis. Mechanism of the					
111	1	22 and CO2 through lungs, arterial and venous circulation. Bohr effect,					
		rbon dioxide binding haemoglobin. pH maintenance by cellular and					
	intracellular proteins. Phosphate and bicarbonate buffers, Metabolic acidosis and alkalosis. Respiratory acidosis and alkalosis. Regulation of fluid and electrolyte						
	balance.						
IV	Sensory transd	uction, Nerve impulse transmission- nerve cells, synapses, reflex arc					
	structure, resting membrane potential, Nernst equation, action potential, voltage gaion-channels, impulse transmission, neurotransmission, neurotransmitter recept						
	synaptosomes, synaptotagmin, rod and cone cells in the retina, changes in the v						
		emical reaction and regulation of rhodopsin, odour receptors, learning					
	=	Chemistry of muscle contraction – actin and myosin filaments, theories					
involved in muscle contraction, mechanism of muscle contraction, energy							
	muscle contract	tion.					

V Reproductive system- sexual differentiation and development; sperm transport, sperm capacitation, semen analyses and Acrosome reaction. Clinical relevance of female reproductive physiology- menstrual cycle, pregnancy and menopause. Fertilisation and infertility issues.

Hormones – Classification, Biosynthesis, circulation in blood, modification and degradation. Mechanism of hormone action, Target cell concept. Hormones of Hypothalamus, pituitary, Pancreatic, thyroid & parathyroid, adrenal and gonadal hormones. Synthesis, secretion, physiological actions and feedback regulation of synthesis.

### **Reading List (Print and online)**

- 1. https://www.genome.gov/genetics-glossary/Cell-Cycle
- 2.https://my.clevelandclinic.org/health/diseases/16083-infertility-causes
- 3. https://www.webmd.com/heartburn-gerd/reflux-disease
- 4. https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5760509/
- 5. https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3249628/

### **Self-Study**

- 1. Variation in cell differentiation and progression
- 2. Lesch Nyhan syndrome, orotic aciduria and GERD

### **Recommended Texts**

- 1. Karp, G. (2010) Cell and Molecular Biology: Concepts and Experiments (6th ed), John Wiley & Sons. Inc.
- 2.Bruce Alberts and Dennis Bray (2013)Essential Cell Biolog(4<sup>th</sup> ed),Garland Science.
- 3. De Robertis, E.D.P. and De Robertis, E.M.F. (2010) Cell and Molecular Biology(8<sup>th</sup> ed), Lippincott Williams and Wilkins, Philadelphia.
- 4.Cooper, G.M. and Hausman, R.E. (2009), The Cell: A Molecular Approach (5<sup>th</sup> ed). Sunderland, Mass. Sinauer Associates, Inc.
- 5. Wayne M. Baker (2008) The World of the Cell. (7<sup>th</sup> ed). Pearson Benjamin Cummings Publishing, San Francisco. Cell Biology.
- 6. John E. Hall (2010) Guyton and Hall Textbook of Medical Physiology (12<sup>th</sup> ed), Saunders.
- 7. Harrison's Endocrinology by J. Larry Jameson Series: Harrison's Specialty, 19th Edition Publisher: McGraw-Hill, Year: 2016.

#### **Method of Evaluation:**

Test I	Test II	Assignment	Assignment Seminar		Total
5	10	5	5	75	100

### **Methods of Assessment:**

Recall (K1) - Simple definitions, MCQ, Recall steps, Concept definitions.

**Understand/ Comprehend (K2) -** MCQ, True/False, Short essays, Concept explanations, Short summary or overview.

Application (K3) - Suggest idea/concept with examples, Solve problems, Observe, Explain.

**Analyse** (**K4**) -Problem-saving questions, Finish a procedure in many steps, Differentiate between various ideas.

Evaluate (K5) - Longer essay/ Evaluation essay, Critique or justify with pros and cons.

Create (K6)- Check knowledge in specific or offbeat situations. Discussion.

### **Mapping with Programme Outcomes:**

	PO 1	PO 2	PO 3	PO 4	PO 5	PO 6	PO 7	PO 8	PO 9	PO 10
CO 1	S	S	S	S	S	M	S	S	S	M
CO 2	S	S	S	S	S	L	S	S	S	M
CO 3	S	S	S	S	S	M	M	S	S	M
CO 4	S	S	S	S	S	M	M	S	S	M
CO 5	M	S	L	S	S	L	M	M	L	L

S - Strong M - Medium L - Low

Course / Course Code	ELECTIVE COURSE - II/ 23P1B4EC					
Title of the Course:	MICROBIOLOGY AND IMMUNOLOGY					
Semester / Credits:	I / 3					
Pre-requisites, if any	The student should possess basic knowledge about					
	microorganisms,types					
	and their general characteristics. The students are also expected to possess					
	basic understanding about the process of infection, immunological					
	defense and pathological outcomes, if any.					
Course Objectives:	<ol> <li>To appreciate the classification of microorganisms based on their structure, size and shape with an insight into the ancient scriptures aboutmicrobes.</li> <li>To understand the role of microorganisms in environment and also</li> </ol>					
	to learn the cultureconditions.  3. To recognize the possible contamination of foods by					
	microorganisms, to learn about counteracting preservative measures					
	and to know about probiotic nature ofmicroorganisms.					
	4. To gain knowledge on pathogenic mediation by microorganisms					
	<ul><li>and preventive measures as well.</li><li>5. To comprehend the features of antimicrobial agents, their</li></ul>					
	mechanism of action along with the side effects and also toexplore					
	natural remedial measures againstmicrobes.					
	6. Tobeabletoexploitthevariousfeaturesofmicroorganismsforthe					
Course Outcomes:	beneficial industrial production.  On successful completion of the course, students should be able to:					
Course Outcomes:	CO1: To classify (by both ancient and modern modes) different types					
	of					
	microorganismsandexplainlifecycleofthemicrobes(K1,K2&K5)					
	CO2: To recognize the microorganisms involved in decay of foods and willbeabletoapplyvariouscounteractingmeasures. The students als o will be able to relate the role of certain beneficial microbes in day-to-day's food consumption (K1, K2 &K4).					
	CO3: To understand the common pathogenic bacteria and fungi that cause toxic effects and also will be able to employ curative measures (K1 & K2).					
	CO4:To analysevarious features of widevariety of antimic robial agents along with their mode of action, in addition, being able to apprehend the valuable potentials of traditional and easily available herbs (K2, K5 & K6).					
	CO5:To applyknowledgegainedinproductionofindustriallyimportant products as both pharmaceutical and nutraceutical (K2, K4 &K5).					
	Units					
I Taxonomical cl	assification - bacteria, viruses (DNA, RNA), algae, fungi and protozoa.					
Distribution and	role of microorganisms in soil, water and air. Charaka's classification					
of microbes, lyt	ic cycle and lysogeny. Types of culture media, isolation of pure culture,					

- growth curve and the measurement of microbial growth.
- II Contamination and spoilage of foods cereals, cereal products, fruits, vegetables, meat, fish, poultry, eggs, milk and milk products. General principles of traditional and modern methods of food preservation Removal or inactivation of microorganisms, boiling, steaming, curing, pasteurization, cold processing, freeze drying, irradiation, vacuum packing, control of oxygen and enzymes. Microbes involved in preparation of fermented foods cheese, yoghurt, curd, pickles, rice pan cake, appam, ragi porridge and bread.
- III Food poisoning- bacterial food poisoning, Salmonella, Clostridium blotulinum (botulism), Staphylococcus aureus, fungal food poisoning aflatoxin, food infection Clostridium, Staphylococcus and Salmonella. Pathogenic microorganisms, E. coli, Pseudomonas, Klebsilla, Streptococcus, Haemophilus, & Mycobacterium, causes, control,prevention,cureandsafety.Foodmicrobiologicalscreening-Real time PCR, ELISA, Aerobic and anaerobic Plate Count, dye reduction method, anaerobic lactic acid bacteria, anaerobic spore formers, Hazard analysis critical control point(HACCP)
- IV Antimicrobial chemotherapy, General characteristics of antimicrobial agents. Mechanism of action sulfonamides, sulphones and PAS. Penicillin, streptomycin-spectraofactivity, mode of administration, mode of action, adverse effects and sensitivity test. Antiviral and antiretroviral agents, Antiviral RNA interference, natural intervention (Natural immunomodulators routinely usedin Indian medicalphilosophy).
- Immune system definition and properties. Cells of the immune system neutrophils, eosinophils, basophils, mast cells, monocytes, macrophages, dendritic cells, natural killer cells, and lymphocytes (B cells and T cells). Lymphoid organs- Primary and Secondary; structure and functions. Antigens and Complement System: definition, properties, antigenicity and immunogenicity, antigenic determinants and haptens. Antigen antibody interactions molecular mechanism of binding. Affinity, avidity, valency, cross reactivity and multivalent binding. Immunoglobulins & Immune Response: Structure, classes and distribution of antibodies. Antibody diversity. Immune system in health & disease, Transplantation immunology graft rejection and HLA antigens. Immunological techniques, Flow cytometry and its applications.

### **Reading List (Print and Online)**

- 1. https://www.ijam.co.in/index.php/ijam/article/view/1326 (Krumi (Microorganisms) in Ayurveda- a critical review)
- 2. Virtual Lectures in Microbiology and Immunology, University of Rochester
- 3. https://www.frontiersin.org/articles/10.3389/fphar.2020.578970/full#h9
- 4. https://www.frontiersin.org/articles/10.3389/fmicb.2018.02151/full
- 5. https://www.ncbi.nlm.nih.gov/pmc/articles/PMC7559905/

### **SelfStudy**

- 1. Microbial infections and gut microbiome with relevance to tridoshas
- 2. Microbial population and pH variations in different dairy products.

### **Recommended Texts**

- 1.Michael J.Pelczar Jr(2001) Microbiology (5th ed), McGraw Hill Education (India) Private Limited.
- 2.Frazier WC, Westhoff DC, Vanitha NM (2010) Food Microbiology (5<sup>th</sup> ed), McGraw Hill Education (India) Private Limited.
- 3. Willey J and Sherwood L (2011)Prescott's Microbiology (8<sup>th</sup> ed), McGraw Hill Education (India).

- 4. Ananthanarayanan, Paniker and Arti Kapil (2013) Textbook of Microbiology (9<sup>th</sup> ed), Orient Black Swan.
- 5. Judy Owen, Jenni Punt Kuby (2013), Immunology (Kindt, Kuby Immunology) (7th ed) W. H. Freeman & Co.
- 6.Brooks GF and Carroll KC (2013) Jawetz Melnick&Adelbergs Medical Microbiology,(26<sup>th</sup> ed), McGraw HillEducation.
- 7. Greenwood D (2012) Medical Microbiology, Elsevier Health.

### **Method of Evaluation:**

Test I	Test II	Assignment	Seminar	End Semester Examination	Total
5	10	5	5	75	100

### **Methods of Assessment:**

**Recall (K1) - Simple definitions, MCQ, Recall steps, Concept definitions.** 

**Understand/ Comprehend (K2) -** MCQ, True/False, Short essays, Concept explanations, Short summary or overview.

**Application (K3) -** Suggest idea/concept with examples, Observe, Explain.

**Analyse (K4)-** Finish procedure in stepwise manner, Differentiation between various ideas, Map knowledge.

**Evaluate (K5)** - Longer essay/ Evaluation essay, Critique or justify with pros and cons.

**Create** (**K6**)- Check knowledge in specific or offbeat situations, Discussion, Debating, Presentation.

### **Mapping with Programme Outcomes:**

	PO 1	PO 2	PO 3	PO 4	PO 5	PO 6	PO 7	PO 8	PO 9	PO 10
CO 1	S	L	S	S	S	S	M	S	S	S
CO 2	S	S	S	S	S	M	L	M	S	S
CO 3	S	M	M	S	M	M	M	M	L	M
CO 4	S	M	M	M	M	M	M	S	S	S
CO 5	S	L	S	S	M	L	L	S	S	S

S - Strong M - Medium L - Low

Course / Course Code:	CORECOURSE - III / 23P2B5
TitleoftheCourse:	ENZYMOLOGY
Semester / Credits:	II / 5
Pre-requisites, if any:	Basic knowledge about catalysis, kinetics and chemical reaction
•	Mechanisms.
Course Objectives:	<ol> <li>Students will be introduced to the theory and practice of enzymology.</li> <li>Mechanisms of catalysis and factors affecting catalysis will be understood</li> <li>The kinetics of enzyme catalyzed reactions in the absence and presence of inhibitors will be studied and the options for applying enzymes and their inhibitors in medicine will be analyzed.</li> </ol>
	4. Students will learn about the applications of enzymes in research, medicine, and industry, which will prepare them for careers in industrial and biomedical research.
	5. The control of metabolic pathways and cellular responses through enzyme regulation will be emphasized.
Course Outcomes:	On successful completionofthecourse, students should be able to: CO1:Describe the catalytic mechanisms employed by enzymes (K1, K2 & K5). CO2:Choose and use the appropriate methods to isolate and purify enzymes and check the purity of the enzyme(K1,K2, K3,K4 &K5). CO3:Analyze enzyme kinetic data graphically, calculate kinetic parameters, determine the mechanism of inhibition by a drug/chemical and analyze options for applying enzymes and their inhibitors in medicine (K1, K2, K3 &K4). CO4:Explain allosterism and cooperativity and differentiate Michaelis- Menten kinetics from sigmoidal kinetics. The role played by enzymes in the regulation of vital cellular processes will be appreciated (K1,
	K2, K5, K6). CO5:Highlight the use of enzymes in industries and biomedicine (K1,K2 & K3).
	Units
I Introduction to	o enzymes and features of catalysis: A short history of the discovery o

	Cints
I	Introduction to enzymes and features of catalysis: A short history of the discovery of enzymes and how they became powerful biochemical tools. Holoenzyme, apoenzyme, cofactors, coenzyme, prosthetic groups, Classification and Nomenclature, Specificity of enzyme action-group specificity, absolute specificity, substrate specificity, stereochemical specificity. Active site, Identification of amino acids at the active site-trapping of ES complex, identification using chemical modification of amino acid side chains and by site-directed mutagenesis.  Mechanisms of enzyme catalysis: acid-base catalysis, covalent catalysis, electrostatic catalysis, metal ion catalysis, proximity and orientation effects, Low barrier H-bonds, Structural flexibility Mechanism of action of chymotrypsin.
II	Enzyme techniques: Isolation and purification of enzymes - Importance of enzyme
	purification, methods of purification- choice of source, extraction, fractionation methods-based on size or mass (centrifugation, gel filtration); based on polarity (ion-exchange chromatography, electrophoresis, isoelectric focusing, hydrophobic

interaction chromatography); based on solubility (change in pH, change in ionic strength); based on specific binding sites (affinity chromatography), choice of methods, Criteria of purity of enzymes. Enzyme units - Katal, IU. Measurement of enzyme activity - discontinuous, continuous, coupled assays; stopped flow method and its applications. Isoenzymes and their separation by electrophoresis with special reference to LDH

Enzyme kinetics I: Thermodynamics of enzyme action, Activation energy, transition-state theory, steady-state kinetics & pre-steady-state kinetics. Single substrate enzyme catalyzed reactions -assumptions, Michaelis-Menten and Briggs-Haldane kinetics, derivation of Michaelis-Menten equation. Double reciprocal (Lineweaver-Burk) and single reciprocal (Eadie-Hofstee) linear plots, their advantages and limitations. Analysis of kinetic data- determination of Km, Vmax, kcat, and their physiological significance, Importance of kcat/Km. Enzyme inhibition: Irreversible inhibition. Reversible inhibition-Competitive, uncompetitive, noncompetitive, mixed and substrate inhibition. Michaelis-Menten equation in the presence of competitive, uncompetitive and noncompetitive inhibitors. Graphical analysis - Diagnostic plots for the determination of inhibition type. Therapeutic use of enzyme inhibitors-Aspirin, statins (irreversible inhibitors), Methotrexate (competitive inhibitor), Etoposide (non-competitive inhibitor), camptothecin (uncompetitive inhibitor).

Demonstration: Using Microsoft Excel to Plot and Analyze Kinetic Data

Enzyme kinetics II: Allosteric enzymes: Cooperativity, MWC and KNF models of allosteric enzymes, Sigmoidal kinetics taking ATCase as an example. Regulation of amount and catalytic activity by - extracellular signal, transcription, stability of mRNA, rate of translation and degradation, compartmentation, pH, temperature, substrate concentration, allosteric effectors, covalent modification. Regulation of glycogen synthase and glycogen phosphorylase. Feedback inhibition-sequential, concerted, cumulative, enzyme-multiplicity with examples.

Bi - Substrate reactions: Single Displacement reactions (SDR) (Ordered and Random bi bi mechanisms), Double Displacement reactions (DDR) (Ping pong mechanism), Examples, Cleland's representation of bisubstrate reactions, Graphical analysis (diagnostic plots) to differentiate SDR from DDR.

Enzyme technology: Immobilization of enzymes — methods - Reversible immobilization (Adsorption, Affinity binding), Irreversible immobilization (Covalent coupling, Entrapment and Microencapsulation, Crosslinking, Advantages and Disadvantages of each method, Properties of immobilized enzymes. Designer enzymes- ribozymes and deoxyribozymes, abzymes, synzymes. Enzymes as therapeutic agents-therapeutic use of asparaginase and streptokinase. Application of enzymes in industry- Industrial application of rennin, lipases, lactases, invertase, pectinases, papain.

### Reading List (Print and Online)

 Enzymes | MIT OpenCourseWare | Free Online Course Materials https://ocw.mit.edu/high-school/biology/exam-prep/chemistry-of-life/enzymes/

### 2. Enzymology

V

https://onlinecourses.swayam2.ac.in/cec20\_bt20/preview https://mooc.es/course/enzymology/

### 3. The active site of enzymes

https://dth.ac.in/medical/courses/biochemistry/block-1/1/index.php

### 4. Enzymes and Enzyme Kinetics

https://www.lecturio.com/medical-courses/enzymes-and-enzyme kinetics.course#/

Mechanistic enzymology in drug discovery: a fresh perspective

https://www.nature.com/articles/nrd.2017.219

Enzyme Biosensors for Biomedical Applications: Strategies for Safeguarding Analytical Performances in Biological Fluids

https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4934206/

### **Self Study**

- 1. Mechanistic enzymology in drug discovery
- 2. Enzyme Biosensors for Biomedical Applications

#### **Recommended Texts**

1. Enzymes: Biochemistry, Biotechnology and Clinical chemistry, 2nd edition, 2007,

Palmer T and Bonner P; Affiliated- East West press private Ltd, New Delhi.

- 2.Fundamentals of Enzymology, 3rd edition, 2003, Price NC and Stevens L; Oxford University Press, New York.
- 3. Voet's Biochemistry, Adapted ed, 2011, Voet, D and Voet JG; Wiley, India
- 4.Lehninger Principles of Biochemistry, 8th edition, 2021, .Nelson DL and Cox MM; WH Freeman & Co, New York.
- 5. Biochemistry, Berg JM, Stryer L, Gatto,G, 8th ed, 2015;WH Freeman & Co., New York.
- 6.Enzyme Kinetics and Mechanism; Cook PF, Cleland W, ;2007; Garland Science, London.

### **Method of Evaluation:**

Test I	Test II	Assignment	Seminar	End Semester Examination	Total
5	10	5	5	75	100

### **Methods of Assessment:**

**Recall (K1) - Simple definitions, MCQ, Recall steps, Concept definitions.** 

**Understand/ Comprehend (K2) -** MCQ, True/False, Short essays, Concept explanations, Short summary or overview.

**Application (K3)** - Suggest idea/concept with examples, Solve problems, Observe, Explain.

**Analyse (K4)-** Problem-solving questions, Finish a procedure in many steps, Differentiate between various ideas.

**Evaluate (K5)** - Longer essay/ Evaluation essay, Critique or justify with pros and cons.

Create (K6)- Check knowledge in specific or offbeat situations, Discussion.

### **Mapping with Programme Outcomes:**

	PO 1	PO 2	PO 3	PO 4	PO 5	PO 6	PO 7	PO 8	PO 9	PO 10
CO 1	S	M	S	L	M	S	L	S	S	M
CO 2	S	S	S	S	M	M	L	S	S	S
CO 3	S	S	S	S	M	M	M	S	S	S
CO 4	S	S	S	S	M	M	M	S	S	S
CO 5	S	S	S	S	M	L	M	S	S	S

S - Strong

M - Medium

L - Low

Course / Course Code:	CORECOURSE - IV / 23P2B6						
TitleoftheCourse:	CELLULAR METABOLISM						
Semester / Credits:	II / 5						
Pre-requisites, if any:	Basic knowledge on biochemical reactions such as addition, deletion,						
	rearrangement, transfer and breaking of bonds.						
Course Objectives:	1. Familiarize on blood glucose homeostasis						
	2. Provide an insight into the metabolic path way of glycogen, glycoprotein, mucopolysaccharide and peptidoglycan with						
	clinical correlation wherever required						
	3. Inculcate knowledge on nucleotide metabolism and disorders						
	associated with it						
	4. Provide a platform to understand the versatile role of PLP in amino acid degradation, formation of specialized products and						
	disorders associated with ammonia detoxification						
	5. Educate on heme and sulphur metabolism with associated clinical						
	manifestation.						
<b>Course Outcomes:</b>	On successful completion of the course, students should be able to:						
	CO1: Appreciate the modes of synthesis and degradation of glucose						
	and will be able to justify the pros and cons of maintain the						
	blood sugar level (K1, K2& K5).						
	CO2: Gain knowledge on polysaccharide metabolism and glycogen						
	storage disease (K1, K2& K5). CO3: Acquaint with the making and braking of nucleotides (K1,						
	K2& K4).						
	CO4: Differentiate the diverse reaction a particular amino acid can						
	experience (K1, K2& K3).						
	CO5: Correlate the disturbance of metabolic reactions to clinical						
	manifestations with reference to heme and sulphur metabolism						
	(K1, K2, K4& K5).						
I Clysolysis se	Units						
	robic and anaerobic, inhibitors and regulation. Feeder pathway- entry						
	to glycolysis, Galactosemia, fructosuria, Pyruvate dehydrogenase						
_	anism and regulation. Glyoxalate cycle and its regulation.						
	is- source, key enzymes, reaction sequence and its regulation. Blood						
	ostasis and the role of hormones. Pentose phosphate pathway-						
	and its regulation. Metabolism of glycogen and its						
regulation.Bios							
mucopolysacch	arides, Chondroitin sulphate.						

II Oxidation of fatty acids-oxidation of saturated and unsaturated fatty acids ( $\alpha$ ,  $\beta$  &  $\omega$ oxidation) Oxidation of fatty acids with odd and even numbered carbon atoms. Regulation of β oxidation. Ketogenesis and its regulation. Biosynthesis of fatty acid and unsaturated, chain elongation, regulation. Biosynthesis of prostaglandins, thromboxanes and leukotrienes and hydroxyl eicosanoic acids. Biosynthesis and degradation of triacylglycerol, phosphoglycerolipids-lecithin, cephalin, plasmalogens and phosphatidyl inositol, Sphingolipid-sphingomyelin, cerebrosides, sulfatides and gangliosides. Cholesterol biosynthesis and its regulation. Lipoprotein metabolism-chylomicrons, VLDL, HDL and LDL. III Metabolism of nucleotides- De novo synthesis and salvage pathways of purine and pyrimidine nucleotides. Regulation and inhibitors of nucleotide biosynthesis. Role of ribonucleotide reductase and its regulation. Degradation of purine and pyrimidine nucleotides. Biosynthesis of non-essential amino acids. Role and biological significance of IVglutamate dehydrogenase, glutamine and asparagine synthetase, lysine, proline and phenylalanine hydroxylase. Interconversion of amino acids - proline to glutamate, methionine to cysteine, serine to glycine. Biosynthesis of spermine and spermidine. Degradation of amino acids -glucogenic and ketogenic amino acids. Formation of acetate from leucine and aromatic amino acids, pyruvate from cysteine, threonine and hydroxy proline, α-keto glutarate from histidine and proline, succinate from methionine, threonine, valine and isoleucine, Oxaloacetate from aspartate, glycine and serine. Biosynthesis and degradation of heme. Jaundice-classification, pathology and  $\mathbf{V}$ Differential diagnosis. Oxidation and reduction of inorganic sulphur compounds by microbes and plants. Sulpho transferases and their biological role-rhodanases, sulphatases, 3-mercapto pyruvate sulphur transferases. Mucopolysaccharidoses -Hunter syndrome, Sanfilippo syndrome and Maroteaux-Lamy syndrome. Oxidation of cysteine to sulphate and inter conversion of sulphur compounds. Reading List (Print and Online) 1. https://www.embopress.org/doi/full/10.1038/msb.2013.19 2. https://people.wou.edu/~guralnl/450Glycogen%20metabolism.pdf 3. https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3243375/ 4. https://www.researchgate.net/publication/334458898 Urea Cycle 5.https://www.researchgate.net/publication/51233381\_Heme\_biosynthesis\_and\_its\_regu lation Towards understanding and improvement of heme biosynthesis in filament ous\_fungi 6.https://www.researchgate.net/publication/349746691\_Microbial\_Sulf ur\_Metabolism\_and\_Environmental\_Implications Selfstudy 1. Cori's Cycle and Glucose-Alanine Cycle 2. Coenzymes involved in Methanogenesis

### **Books Recommended**

- 1.David L.Nelson and Michael M.Cox (2012) Lehninger Principles of Biochemistry (6th ed), W.H.Freeman
- 2. Voet.D and Voet. J.G (2010) Biochemistry, (4th ed), John Wiley & Sons, Inc.
- 3.Metzler D.E (2003). The chemical reactions of living cells (2nd ed), Academic Press.
- 4. Zubay G.L (1999) Biochemistry, (4th ed), Mc Grew-Hill.
- 5. Textbook of Biochemistry with Clinical Correlations, 7th Edition, Thomas M. Devlin (Editor), Wiley
- 6. Human Biochemistry James M.Orten & Otto.W.Neuhan- 10th edn- The C.V.Mosby Company.

### **Method of Evaluation:**

Test I	Test II	Assignment	Seminar	End Semester Examination	Total
5	10	5	5	75	100

### **Methods of Assessment:**

**Recall (K1) - Simple definitions, MCQ, Recall steps, Concept definitions.** 

**Understand/ Comprehend (K2) -** MCQ, True/False, Short essays, Concept explanations, Short summary or overview.

**Application** (**K3**) - Suggest idea/concept with examples, Solve problems, Observe, Explain.

**Analyse** (**K4**)- Problem-solving questions, Finish a procedure in many steps, Differentiate between various ideas.

**Evaluate (K5)** - Longer essay/ Evaluation essay, Critique or justify with pros and cons.

Create (K6)- Check knowledge in specific or offbeat situations, Discussion.

### **Mapping with Programme Outcomes:**

	PO 1	PO 2	PO 3	PO 4	PO 5	PO 6	PO 7	PO 8	PO 9	PO 10
CO 1	S	M	S	M	S	M	S	S	S	M
CO 2	S	M	S	S	S	M	S	S	S	M
CO 3	S	M	S	S	S	M	S	S	S	S
CO 4	S	M	S	M	S	M	S	S	S	M
CO 5	S	M	S	S	S	M	S	S	S	S

S-Strong M-Medium L-Low

COREPRACTICAL - II / 23P2BP2
LABORATORY COURSE ON ENZYMOLOGY, MICROBIOLOGY AND CELL BIOLOGY
II / 4
Knowledge on basic principles, Instrumentation of Biochemical techniques and metabolic reactions.
<ol> <li>To inculcate skill in students enabling them to apprehend the widerknowledgeaboutprinciplesandtechniquestobeemployedforthe assayofenzymesunderinvestigation.</li> <li>Toinculcatetheknowledgeofisolationandpurificationtechniquesofen zymes using alkalinephosphataseas an example</li> <li>Toperformexperimentstostudythefactors affectingenzymeactivity</li> <li>Toachievetraininginassayofenzymes</li> <li>Toachievetraininginbasicmicrobiologicaltechniques—preparationofculture,sterilization and staining methods.</li> <li>Toperformthebloodgroupingtestandtopreparebloodsmeartostudydif ferenttypesof blood cells</li> <li>Tolearn molecularbiologytechniqueslikeGelelectrophoresisandBlottingtech niques</li> </ol>
8. Tointroduceindustrialvisitsothatstudentsmaybeawareofactualneedo
On successful completionofthecourse, students should be able to: CO1:  Thestudentwillbeabletoemploytherelevanttechniquesforisolatio nand purificationofenzymes and gainskill inkineticstudies which is essential forresearch activity (K1, K2&K4). CO2:Student will acquire ability in performing enzymeassay, and explicate the methods that form the basis of enzyme characterization (K1, K2& K4). CO3:Learn the Basic concepts in microbiology and cell biology which will be helpful for interdisciplinary research work (K1, K3& K4). CO4: Students will be trained in separation techniques used in molecular Biology which will be supportive in their futureresearch(K1, K3, K4 &K6). CO5: Industrial visits will provide the students with an opportunity to learn practically through interaction, working methods and employment practices. Students will have an exposure to Industrial standard and current work practices(K1,K2,K3,K4 &K6). Units

### Enzymology Demonstration Preparation of various pH buffer solutions using pH meter Determination of activity Effect of pH Effect of Temperature Determination of Km by Line Weaver – Burk Plot Acid phosphatase Determination of activity Effect of pH Effect of Temperature Determination of specific activity Determination of Km by Line Weaver – Burk Plot Alkaline phosphatase Determination of activity Effect of pH Effect of Temperature Determination of specific activity Determination of Km by Line Weaver – Burk Plot Urease Determination of activity Effect of pH Effect of Temperature Determination of specific activity Determination of Km by Line Weaver – Burk Plot Microbiology: Sterilization, Culture and inoculum preparation II **Physiology & Cell Biology**: Test for blood grouping (Haemagglutination) Ш Group Experiments: Separation of proteins based on molecular weight by SDS PAGE IVIndustrial visit can be organized to students through Academia–Industry collaborative Program **Reading List (Print and Online)** 1.https://www.researchgate.net/publication/337146254\_Kinetic\_studies\_with\_alkaline\_p hosphatase 2.https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4846332/ 3.https://www.ijsr.net/archive/v3i8/MDIwMTU0MDk=.pdf 4.https://www.researchgate.net/publication/349318898\_ABC\_of\_ Periheral\_smear 5.https://ncdc.gov.in/WriteReadData/1892s/File608.pdf 6.https://www.ncbi.nlm.nih.gov/books/NBK562156/ SelfStudy 1. Safety measures and Good Laboratory Practices in microbiology laboratory 2. Staining of bacteria – Gram Staining 3. Peripheral Blood smear – Staining and Interpretation 4. Agarose gel electrophoresis of genomic DNA

#### **Books Recommended**

- 1. David Plummer (2001) An Introduction to Practical Biochemistry (3rd ed) McGraw Hill Education (India) Private Ltd.
- 2. Jayaraman, J (2011) Laboratory Manual in Biochemistry, New age publishers.
- 3. Fundamentals of Enzymology; 3rd Edn. Nicholas C. Price and Lewis Stevens, Oxford University Press (2012).
- 4. Enzymes: A Practical Introduction to Structure, Mechanism, and Data Analysis; Robert A. Copeland, Wiley-VCH Publishers (2000).
- 5. Cappuccino JG & Sherman N (2005) Microbiology-A Laboratory Manual, Pearson Education Inc.
- 6. Practical Enzymology, Second Revised Edition: Hans Bisswanger, Wiley Blackwell; 2 edition (2011).

### **Method of Evaluation:**

Test I	Test II	End Semester Examination	Total	Grade
20	20	60	100	

### **Methods of Assessment:**

**Recall (K1) - Simple definitions, MCQ, Recall steps, Concept definitions.** 

**Understand/ Comprehend (K2) -** MCQ, True/False, Short essays, Concept explanations, Short summary or overview.

**Application (K3) -** Suggest idea/concept with examples, Solve problems, Observe, Explain.

**Analyse (K4) -** Problem-solving questions, Finish a procedure in many steps, Differentiate between various ideas.

**Evaluate (K5)** - Longer essay/ Evaluation essay, Critique or justify with pros and cons.

Create (K6) - Check knowledge in specific or offbeat situations, Discussion.

### **Mapping with Programme Outcomes:**

	PO 1	PO 2	PO 3	PO 4	PO 5	PO 6	PO 7	PO 8	PO 9	PO 10
CO 1	S	S	S	S	M	S	L	S	M	S
CO 2	S	S	S	S	M	S	L	S	M	S
CO 3	S	S	S	S	M	S	M	S	M	S
CO 4	S	S	S	S	S	S	S	S	S	S
CO 5	S	S	S	S	S	S	S	S	S	S

S - Strong M - Medium

L-Low

Course / Course Code:	ELECTIVE COURSE - III / 23P2B7EC
Title of the Course:	MOLECULAR BIOLOGY
Semester / Credits:	П/3
Pre-requisites, if any:	Knowledge of the basics of genetics, cell biology and molecular
	biology.
Course Objectives:	1. To introduce the students to the process of inheritance, concepts of
	genes, genome, chromatin and chromosomes.
	2. To impart a thorough understanding of the key events of molecular
	biology, including the mechanisms of DNA replication,
	transcription and translation along with DNA repair mechanisms.  3. To provide a detailed understanding of post transcriptional and
	posttranslational modifications and processing of eukaryotic
	RNA and proteins.
	4. To give a detailed explanation of transcriptional regulation with
	lac operon and tryptophan operon as examples.
	5. To impart adequate information of the types of regulatory RNAs
C Ot	along with key concepts of gene silencing.
Course Outcomes:	On successful completion of the course, students should be able to: CO1: Comprehend the organization of genomes, the molecular basis
	of DNA replication, recombination and transposition, the
	significance of these processes, the various ways in which the
	DNA can be damaged leading to mutations and lesions and the
	different ways in which they are repaired (K1, K2, K3& K5).
	CO2: Gain knowledge about how genes are transcribed and translated
	in prokaryotes and eukaryotes and how these processes are
	regulated, recognize the nature of the genetic code and the various experimental approaches used to crack the code (K1,
	K2, K3, K4& K5).
	CO3: Acquire knowledge of the molecular basis of RNA processing
	and RNA splicing and the various human pathologies that can
	result from defects of RNA modification (K1, K2, K4& K5).
	CO4: Comprehend the techniques of gene silencing and its
	applications (K1, K2, K3, K4, K5& K6).
	CO5: Apply the knowledge they have gained in understanding the
	above vital life processes to enhancing their analytical and problem-solving skills and develop an interest to pursue high
	quality research (K2, K3, K4, K5& K6).
	Units
I Mendel's laws	of inheritance-dominance-complete, incomplete and co- dominance,
	gene mapping in haploids and diploids, recombination mapping-
restriction mapp	ing- modes of gene information transfer in bacterial- conjugation,
	nd transduction. The bacterial chromosome, the eukaryotic genome-
	ructure – Histones, Nucleosome, chromatin- heterochromatin,
	omatin remodeling, DNAase hypersensitive sites, genome organization –
	adox, reassociation kinetics, repetitive sequences, gene amplification, ogenes, split genes, organelle genomes – mitochondrial and chloroplast
teromeres, pseud	zenes, spin genes, organene genomes – initoenonariai ana emoropiast

	genome.
II	DNA replication and repair: Enzymes of replication, prokaryotic replication mechanisms, primosome & replisomes, eukaryotic DNA replication, the role of topoisomerases and telomerase, regulation of replication, difference between prokaryotic and eukaryotic replication. Mutations -Types of mutations, mechanisms of mutations, mutagenic agents. DNA repair mechanisms – Direct repair, excision repair, mismatch repair, recombination repair, SOS response, eukaryotic repair systems. Recombination and mobile genetic elements- the Holliday model, the general recombination in <i>E.coli</i> , site specific
III	recombination, transposons and retroposons.  Transcription – Prokaryotic transcription-subunits of RNA polymerase, E. coli promoters,
	sigma factor and promoter recognition, alternative sigma factors, initiation, elongation, Rho-dependent and independent termination of transcription. Eukaryotic transcription-Initiation, promoter elements, RNA polymerases, transcription factors, regulatory sequences in eukaryotic protein – coding genes, CpG islands, enhancers.  Translation – organization of the ribosome, the genetic code, evidence for a triplet code, deciphering the genetic code, wobble hypothesis, deviation in the genetic code, unusual codons. activation, initiation, elongation and termination of translation in E. coli. The role of tRNA and rRNA, suppressor tRNAs and inhibitors of protein synthesis. Comparison of
	prokaryotic translation with eukaryotic translation.
IV	Regulation of gene expression in prokaryotes—Positive and negative control, the lac operon, identification of operator and regulator sequences by mutations, induction and repression, Foot-printing and gel-shift assays for identification of protein-DNA interactions. Catabolite repression. <i>Trp</i> operon — Attenuation, alternative secondary structures of <i>trp</i> mRNA.  Regulation of gene expression in eukaryotes—Response elements, DNA-binding motifs, steroid receptors, association of methylation and histone acetylation with gene expression.
V	Post transcriptional modifications in eukaryotes- RNA processing- mRNA 5' capping and
	3'poly-adenylation, introns and exons, RNA splicing,- spliceosome assembly, alternative splicing, processing of tRNA and rRNA, self-splicing, ribozymes, RNA editing-substitution and insertion/deletion editing, Genome editing-CRISPR- Cas technology Post translational modification of proteins- Proteolytic cleavage, covalent modifications, glycosylation of proteins, disulfide bond formation, Protein sorting – signal peptides, transport of secretory proteins, Golgi and post-golgi sorting, coated vesicles, targeting of mitochondrial, lysosomal and nuclear proteins, Protein degradation-Ubiquitination of proteins, Protein folding-chaperones
	Reading List (Print and Online)
	<ol> <li>Molecular Biology Free Online Course by MIT Part 3: RNA Uploaded by edX</li> <li>https://mooc.es/course/molecular-biology/</li> </ol>
	3. https://onlinecourses.swayam2.ac.in/cec20_ma13/preview
	4. https://learn.genetics.utah.edu/
	5. https://www.cellbio.com/education.html
	6. https://lifescienceinteractive.com/category/molecular-biology/
	Self Study  1. Multiple roles of poncoding PNAs (long ncPNA siPNA miPNA) in
	1. Multiple roles of noncoding RNAs (long ncRNA, siRNA, miRNA) in development and differentiation; implication of ncRNAs in pathologies.
	development and differentiation, implication of next vas in pathologies.

2. mRNA degradation-nonsense-mediated decay.

### **Recommended Texts**

- 1. Lewin's Genes XII: 12th edition, Krebs JE, Goldstein ES, Kilpatrick ST; Prentice Hall, Delhi.
- 2. Molecular Biology of the Gene: 6th edition, Watson JD, Baker TA, Bell S, Gann A, Levine M, Losick R; Cold Spring Harbor Laboratory Press, New York.
- 3.Essential Cell Biology:3rd edition, Alberts B, Bray D, Hopkin K, Johnson A, Lewis J, Raff M, Roberts K, Walter P; Garland Science, New York.
- 4. Molecular Cell Biology: 8th edition, Lodish H, Arnold Berk; W.H.Freeman & Co, New York.
- 5. Karp's Cell and Molecular Biology: Concepts and Experiments, 8th Edition; Wiley, India.
- 6. An Introduction to Genetic Analysis 12th edition, Griffith A. F, Doebley J, Peichel C, David A, Wassarman DA; Albion Press.W.H.Freeman & Co,New York.

#### **Method of Evaluation:**

Test I	Test II	Assignment	Seminar	End Semester Examination	Total
5	10	5	5	75	100

#### **Methods of Assessment:**

**Recall (K1) - Simple definitions, MCQ, Recall steps, Concept definitions.** 

**Understand/ Comprehend (K2) -** MCQ, True/False, Short essays, Concept explanations, Short summary or overview.

**Application (K3) -** Suggest idea/concept with examples, Solve problems, Observe, Explain.

**Analyse (K4)-** Problem-solving questions, Finish a procedure in many steps, Differentiate between various ideas.

**Evaluate (K5)** - Longer essay/ Evaluation essay, Critique or justify with pros and cons.

Create (K6)- Check knowledge in specific or offbeat situations, Discussion.

### **Mapping with Programme Outcomes:**

	PO 1	PO 2	PO 3	PO 4	PO 5	PO 6	PO 7	PO 8	PO 9	PO 10
CO 1	S	S	S	L	M	L	S	S	S	S
CO 2	S	S	S	M	M	L	M	S	S	S
CO 3	S	S	S	L	M	L	M	S	S	S
CO 4	S	S	S	M	M	L	S	S	S	S
CO 5	S	S	S	S	S	M	M	S	S	S

S - Strong M - Medium L - Low

Course	e / Course Code:	ELECTIVE COURSE - IV / 23P2B8EC				
Titled	oftheCourse:	ENERGY AND DRUG METABOLISM				
Semest	ter / Credits:	II/3				
Pre-re	equisites, If any:	Basic knowledge on biochemical reactions such as addition, deletion, rearrangement, transfer and breaking of bonds.				
Cour	se Objectives:	<ol> <li>Familiarize on concepts of enthalpy, entropy, free energy, redox system, biological oxidation and high energy compounds.</li> <li>Provide an insight into the relationship between electron flow and phosphorylation.</li> <li>Inculcate knowledge on processes involved in converting light energy to chemical energy and associated food production by autotrophs.</li> <li>Provide a platform to understand the versatile role of Krebs cycle, transport of NADH across mitochondrial membrane and</li> </ol>				
		<ul><li>energetic.</li><li>5. Educate on the various phases xenobiotic metabolism.</li></ul>				
Course Outcomes:		<ul> <li>On successful completionofthecourse, students should be able to:</li> <li>CO1: Appreciate the relationship between free energy and redox potential and will be able to justify the role of biological oxidation and energy rich compounds in maintaining the energy level of the system (K1, K2, K3&amp; K4).</li> <li>CO2: Gain knowledge on role of mitochondria in the production of energy currency of the cell (K1, K2, K5&amp; K6).</li> <li>CO3: Acquaint with the process of photosynthesis (K1, K2&amp; K5).</li> <li>CO4: Comprehend on the diverse role of TCA cycle and the energy obtained on complete oxidation of glucose and fatty acid (K1, K2, K4&amp; K5).</li> <li>CO5: Correlate the avenues available to metabolize the xenobiotics (K1, K2, K4&amp; K5).</li> </ul>				
		Units				
I	Thermodynamic- principles in biology- Concept of entropy, enthalpy and free energy change. Redox systems. Redox potential and calculation of free energy. Biological oxidation — Oxidases, dehydrogenases, hydroperoxidases, oxygenases. Energy rich compounds — phosphorylated and non-phosphorylated. High energy linkages.					
II	Electron transport chain-various complexes of ETC, Q-cycle. Inhibitors of ETC. Oxidative phosphorylation-P/O ratio, chemiosmotic theory. Mechanism of ATP synthesis - role of F0-F1 ATPase, ATP-ADP cycle. Inhibitors of oxidative phosphorylation, ionophores, protonophores. Regulation of oxidative phosphorylation.					
III	Light reaction-Hills reaction, absorption of light, photochemical event. Photo ETC-cyclic and non-cyclic electron flow. Photophosphorylation-role of CF0-CF1 ATPase. Dark reaction- Calvin cycle, control of C3 pathway, and Hatch-Slack pathway (C4 pathway), Photorespiration. Synthesis and degradation of starch.					
IV	Interconversion	of major food stuffs. Energy sources of brain, muscle, liver, kidneyand				

	adipose tissue. Amphibolic nature of Citric acid cycle. Anaplerotic reaction. Krebs
	cycle, Inhibitors and regulation of TCA cycle. Transport of extra mitochondrial NADH
	- Glycerophosphate shuttle, malate aspartate shuttle. Energetics of metabolic pathways
	– glycolysis, (aerobic and anaerobic), citric acid cycle, beta oxidation.
V	Activation of sulphate ions – PAPS, APS, SAM and their biological role. Metabolism
	of xenobiotics – Phase I reactions – hydroxylation, oxidation and reduction. Phase II
	reactions – glucuronidation, sulphation, glutathione conjugation, acetylation and
	methylation. Mode of action and factors affecting the activities of xenobiotic enzymes.
	Reading List (Print and Online)
	1.https://chemed.chem.purdue.edu/genchem/topicreview/bp/ch21/gibbs.php
	2.https://www.ncbi.nlm.nih.gov/pmc/articles/PMC7767752/#:~:text=The%20mitochon
	drial%20electron%20transport%20chain,cellular%20ATP%20through%20oxidative
	%20phosphorylation.
	3.https://www.researchgate.net/figure/Oxidative-phosphorylation-in-mitochondrial-
	electron-transport-chain-ETC-and-proton_fig1_230798915
	4.https://www.lyndhurstschools.net/userfiles/84/Classes/851/photosynthesis%20light%
	20&%20dark%20reactions%20ppt.pdf?id=560837
	5.https://bajan.files.wordpress.com/2010/05/amphibolic-nature-of-krebs-cycle.pdf
	6.https://www.sciencedirect.com/topics/medicine-and-dentistry/xenobiotic-
	metabolism#:~:text=Xenobiotic%20metabolism
	%20can%20be%20defined,more%20readily%20excreted%20hydrophilic%20metabo
	lites
	SelfStudy
	1. Calculation of Keq and △G
	2. Interrelationship of carbohydrate, protein and fat metabolism-role of acetyl CoA
	Recommended Texts
	1.David L.Nelson and Michael M.Cox (2012) Lehninger Principles of Biochemistry
	(6th ed), W.H.Freeman
	2. Robert K. Murray, Darryl K. Granner, Peter A. Mayes, and Victor W. Rodwell
	(2012) Harper's Illustrated Biochemistry (29th ed), McGraw-Hill Medical
	3. Metzler D.E (2003) The chemical reactions of living cells (2nd ed), Academic Press.
	4. Zubay G.L (1999) Biochemistry, (4th ed), Mc Grew-Hill.
	5. Devlin RM (1983) Plant Physiology (4th ed), PWS publishers.

# **Method of Evaluation:**

Test I	Test II	Assignment	Seminar	End Semester Examination	Total
5	10	5	5	75	100

 $6. Taiz\ L,$  Zeiger E (2010), Plant Physiology (5th ed), Sinauer Associates, Inc.

### **Methods of Assessment:**

Recall (K1) - Simple definitions, MCQ, Recall steps, Concept definitions.

**Understand/ Comprehend (K2) -** MCQ, True/False, Short essays, Concept explanations, Short summary or overview.

**Application (K3) -** Suggest idea/concept with examples, Solve problems, Observe, Explain.

**Analyse** (**K4**)- Problem-solving questions, Finish a procedure in many steps, Differentiate between various ideas.

Evaluate (K5) - Longer essay/ Evaluation essay, Critique or justify with pros and cons.

Create (K6)- Check knowledge in specific or offbeat situations, Discussion.

### **Mapping with Programme Outcomes:**

	PO 1	PO 2	PO 3	PO 4	PO 5	PO 6	PO 7	PO 8	PO 9	PO 10
CO 1	S	S	S	M	S	M	S	S	S	M
CO 2	S	S	S	S	S	S	S	S	S	S
CO 3	S	S	S	S	S	S	S	S	S	S
CO 4	S	M	S	M	S	M	S	S	S	L
CO 5	S	M	S	S	S	M	S	S	S	S

S - Strong M - Medium L - Low

Course	e / Course Code:	SKILL ENHANCEMENT COURSE [SEC] – I/ 23P2B9SEC				
Title	of theCourse:	NUTRITIONAL BIOCHEMISTRY				
Seme	ester / Credits:	II / 2				
Pre-re	equisites, if any:	Basic knowledge on food, nutrition & dietetics, and metabolism of nutrients.				
Cour	se Objectives:	<ol> <li>To understand basic concepts involved in growth, health, nutrition, physiology and metabolism.</li> <li>To discuss the concepts and applications of nutrition in correlation with biochemistry.</li> <li>To define nutritional needs in healthy individuals and modification of diet during illness.</li> </ol>				
Cour	rse Outcomes:	<ul> <li>On successful completionofthecourse, students should be able to:</li> <li>CO1: Plan a balanced diet based on an individual's energy requirement, Assess nutritional status of an individual(K3, K4&amp; K5).</li> <li>CO2:Describe the biochemical, physiological and nutritional functions of macronutrients and their integrated role.  Understand the role played by antinutritional factors(K1 to K6).</li> <li>CO3:Evaluate the functions of vitamins and minerals ,and fluids and electrolyte balance in different physiological states and in sports persons(K1 to K6).</li> <li>CO4:Identify nutritional deficiency conditions , its prevention and dietary management(K3,K4).</li> <li>CO5: Acquire knowledge about the importance of balanced diet and diet therapy (K5,K6).</li> </ul>				
I	Basic concepts - Nutrition - Food groups and balanced diet. Novel Foods. Calorific value of foods: Direct and indirect calorimetry. Empty calories. Basal metabolic rate: Factors affecting BMR. SDA and physical activity. Calculation of day's energy requirement. Assessment of nutritional status. Lactose intolerance. Nutritional requirement and biochemical changes in different physiological states-infancy, childhood, pregnancy, lactation, and ageing. Sports nutrition.					
II	Elements of nutrition - Plant and animal sources of simple and complex carbohydrates, fats and proteins and their requirement. Biological significance, deficiency and toxicity of macronutrients and micronutrients. Role of dietary fibre. Protein sparing action of carbohydrates and fats. Essential amino acids. Essential fatty acids. Effects of naturally occurring food toxins, preservatives, additives, alcohol and tobacco on health.					
III	Vitamins and Minerals- Dietary sources, classification, biochemical functions, requirements, absorption, metabolism and excretion. Vitamin B complex as coenzyme. Nutritional significance of dietary calcium, phosphorus, magnesium, iron, iodine, zincand copper.					

Malnutrition - Diseases arising due to Protein - Calorie Malnutrition and undernutrition (Kwashiorkor and Marasmus), Prevention of malnutrition. Deficiency diseases associated with vitamin B complex, vitamin C and A, D, E & K vitamins - Mineral deficiency diseases - aetiology, sign and symptoms and dietary supplementation. Enrichment and fortification (vitamins and minerals).  Nutrition in diseases - Aetiology, signs and symptoms, treatment and dietary					
management during fever(Typhoid and Malaria) and infectious diseases(COVID-19),					
Jaundice, hyper acidity (Ulcer), Atherosclerosis, Hypertension, kidney diseases and					
diabetes in adults. Starvation and Obesity. Inter-relationship of nutrition, infection, immunity and poverty.					
Reading List (Print and Online)					
1. https://www.jmedscindmc.com/article.asp?issn=1011-					
4564;year=2014;volume=34;issue=5;spage=211;epage=213;aulast=Shrivastava					
2. https://www.researchgate.net/figure/Relationship-between-malnutrition-infection-and-immunity-Malnutrition-is-considered-the_fig1_280722727					
3. https://en.wikipedia.org/wiki/Novel_food					
4. https://www.chemicalsafetyfacts.org/preservatives/					
5. https://www.sciencedirect.com/topics/agricultural-and-biological-sciences/food-enrichment.					
SelfStudy					
1. Antabuse drugs and food					
2. Selection of foods and market visit, reading and understanding the food labels					
Recommended Texts					
1. Srilakshmi. E(2016) Nutrition Science, New Age International Publishers.					
2. Mahan, Kathleen L (2004) Krause's Food, Nutrition and Diet Therapy,					
W.B.Saunder's 11th Edition.					
3. Andreas M. Papas (1998). Antioxidant Status, Diet, Nutrition, and Health (1st ed) CRC Press.					
4. M. Swaminathan (1995) Principles of Nutrition and Dietetics. Bappeo					
5. Margaret Mc Williams (2012) Food Fundamentals (10th ed), Prentice Hall.					
6. Tom Brody (1998) Nutritional Biochemistry (2nd ed), Academic Press, USA.					

### **Method of Evaluation:**

Test I	Test II	Assignment	ssignment Seminar End Se Exami		Total
5	10	5	5	75	100

### **Methods of Assessment:**

 $Recall (K1) \hbox{-} Simple definitions, MCQ, Recall steps, Concept definitions. \\$ 

 $\label{lem:comprehend} \textbf{Understand/Comprehend(K2)-} MCQ, True/False, Shortessays, Concept explanations, Shortsummary or overview.$ 

 $Application (K3) \hbox{-} Suggestide a/concept with examples, Solve problems, Observe, Explain.$ 

**Analyse** (**K4**) – Problem-saving questions, Finish a procedure in many steps, Differentiate between various ideas.

 $\label{eq:condition} \textbf{Evaluate}(\textbf{K5})\textbf{-} \textbf{Longer essay/ Evaluationessay,} \textbf{Critique or justify with prosand cons.} \\ \textbf{Create}(\textbf{K6})\textbf{-} \textbf{Check knowledge in specific or off beat situations.} \\ \textbf{Discussion.} \\$ 

# **Mapping with Programme Outcomes:**

	PO 1	PO 2	PO 3	PO 4	PO 5	PO 6	PO 7	PO 8	PO 9	PO 10
CO 1	S	S	S	S	S	S	S	S	M	M
CO 2	S	S	S	S	S	S	S	S	M	M
CO 3	S	S	S	S	S	S	S	S	M	M
CO 4	S	S	S	S	S	S	S	S	M	L
CO 5	S	S	S	S	S	S	S	S	M	M

S - Strong M - Medium L - Low

Course / Course Code:	CORE COURSE – V / 23P3B10						
Title of the Course:	CLINICAL BIOCHEMISTRY						
Semester / Credits:	III / 5						
Pre-requisites, if any:	The student should have a basic knowledge of body fluids and their						
	composition and metabolism; anatomy and physiology of vital organs.						
Course Objectives:	1. To understand the need and methods of various biological sample collection.						
	2. To explicitly understand the etiopathogenesis, symptoms and						
	complications of metabolic and hormonal disorders and the relevant						
	diagnostic markers.						
	3. To emphasize the diagnostic significance of serum enzymes in						
	different pathologies and other Laboratory investigations of						
	diagnostic importance so as to differentiate normal from disease.						
	4. To conceive the role of inherited genes in inborn errors of metabolism						
	and methodologies pertaining to <i>in utero</i> diagnosis and post-natal screening.						
	5. To get updated about electrolyte and hormonal imbalances and the						
	biochemical tests to diagnose them.						
Course Outcomes:	CO1: To appreciate the biological significance of sample collection and						
	awareness of the diagnostic/screening tests to detect common						
	non-communicable diseases so as to understand role of						
	laboratory investigations for biochemical parameters and						
	understand the disorders associated with blood cells.						
	CO2: To understand the etiology of metabolic diseases like diabetes and						
	atherosclerosis and avoid such lifestyle disorders by healthy						
	eating and correlate the symptoms with underlying pathology						
	based on diagnostic and prognostic markers.						
	CO3: To understand the diagnostic application of serum/plasma						
	enzymes to correlate their levels with the organ pathologies						
	associated with specific diseases.						
	CO4: To appreciate the role of pre and post-natal diagnosis leading to						
	healthy progeny.						
	CO5: To link the serum hormone levels and clinical symptoms with						
	underlying hormonal disturbances. To review the onward transmission of signal via downstream signaling molecules from						
	cell surface to the nucleus by different pathways by comparing						
	and contrasting them and critically evaluate the network between						
	them resulting in the biological outcome.						
	Units						
	C ARAVU						

Biochemical investigations in diagnosis, prognosis, monitoring, screening: Specimen collection – blood, (primary/Secondary specimen)., urine and CSF. Preservation of biological specimens -blood, urine, CSF and amniotic fluid, Biological reference ranges.

**Disorders of blood cells**: Hemolytic, iron deficiency and aplasticanemia and diagnosis, sickle cell anaemia, thalassemia HBA1C variants. Porphyrias, Thrombocytopenia, Causes of leucopenia, leukemia and leucocytosis. Disorders of blood clotting mechanism - Von willebrand's disease, Hemophilia A, B and C, diagnostic test for clotting disorders,

D-dimer and its clinical significance

Diabetes mellitus: pathology and complications: Acute changes; Chronic complications: Diabetic nephropathy, neuropathy, retinopathy and Diabetic foot ulcers, Random/Fasting/PP glucose testing, Impaired glucose tolerance (IGT), Impaired fasting glucose (IFT), Diagnosis-by GTT, Pre-diabetes, Gestational DM,Glycosylated Haemoglobin (HBA1c); Glycated albumin., Hypoglycaemia and critical alert value for glucose. Markers of complications of Diabetes mellitus: Metabolic syndrome, Lipid profile &lipoproteinemia, Atherosclerosis, Diabetic nephropathy, Micralbuminuira, eGFR.

Point of care testing for glucose (Glucometers) and continuous glucose monitoring (CGM): principle and its use. Major groups of anti-diabetic drugs. Diet and life style modifications.

**III Diagnostic Enzymology:** Clinically Important Enzymes and Isoezyme as diagnostic markers: Clinical significance of AST, ALT, ALP, ACP, CK, γ-GT, amylase, pseudocholinesterase and their pattern in .Myocardial infarction; Liver disease, Bone disease, Muscle disease, Cancer (tumor markers), GI tract pancreatitis); Enzymes as therapeutic agents.

**Pre- and post-natal testing:** Amniocentesis, prenatal detection of inborn errors of metabolism in developing fetus- Autosomal recessive mode of inheritance- cystic fibrosis, X linked recessive inheritance-Duchenne muscular dystrophy. New born screening (NBS) for In born errors of metabolism, Tandem mass spectrometry application in NBS.

**IV Liver function tests:** Liver function test panel, Fatty liver. Plasma protein changes in liver diseases. Hepatitis A,B and C. Cirrhosis and fibrosis. Portal hypertension and hepatic coma. Acute phase proteins -CRP, Haptoglobins, α-fetoprotein, ferritin and transferrin and their clinical significance, Interpreting serum protein electrophoresis. Inflammatory markers (cytokines such as TNF-alpha IL6 and others)

**V** Renal function tests - tests for glomerular and tubular function-Acute and chronic renal failure-Glomerulonephritis, Nephrotic syndrome, uraemia-urinary calculi-Nephrocalcinosis and Nephrolithiasis-causes, pathology and symptoms. Chronic kidney disease. Dialysis-Hemodialysis and peritoneal dialysis.

**Electrolyte disorder:** calcium: hypercalcemia and hypocalcemia; Calcium homoestasis in Blood; phosphate: hyperphosphatemia or hypophosphatemia;

Clinical significance: Potassium: hyperkalaemia and hypokalaemia, Sodium: hypernatremia and hyponatremia; Chloride: hyperchloremia, hyporchloremia

**Hormonal disorders and diagnostics:** T3, T4 and TSH in the diagnosis of thyroid disorders; Diagnostic methods for disorders associated with adrenal, pituitary and sex hormones - Addison's disease, Cushing's syndrome, pituitary tumour, Hypopituitarism, Hypogonadism

# **Reading List (Print and Online)**

## 1. Utility of HIL in Clinical Chemistry:

https://www.aacc.org/science-and-research/clinical-chemistry-trainee-council/trainee-council-in-english/pearls-of-laboratory-medicine/2018/utility-of-hil-in-clinical-chemistry

## 2. Pre, Post and Analytical Errors in Clinical Chemistry laboratory

DOI: 10.7860/NJLM/2016/22587:2173

https://doi.org/10.2147/JMDH.S286679

# 3. Standards of Medical Care in Diabetes—2022 Abridged for Primary Care Providers

https://diabetesjournals.org/clinical/article/40/1/10/139035/Standards-of-Medical-Care-in-Diabetes-2022

https://doi.org/10.2337/diaspect.16.1.32

http://www.ngsp.org/

4. Quality control in clinical laboratory

 $https://www.researchgate.net/publication/335830829\_Quality\_Control\_in\_a\_Clinical\_La~boratory$ 

https://labpedia.net/quality-control-of-the-clinical-laboratory/

https://journals.sagepub.com/doi/full/10.1016/j.jala.2008.12.001

https://doi.org/10.1016/B978-0-12-407821-5.00004-8

https://www.westgard.com/clia.htm

https://www.labroots.com/webinar/bio-rad-unity-solution-molecular-quality-control-data-management

## Self study

## 1. Potential sources of variability in the estimation of the analytes:

Pre-analytical phase: acceptance rejection criteria in terms of haemolysis/icteric/lipemia (HIL) interferences.

Analytical phase: Linearity, detection limits precision, accuracy, specificity, sensitivity; Total Allowable Error. (Definitions and examples).

Post-analytical phase: Units of reporting of clinical chemistry parameters.

# 2. Interpretation of results in clinical chemistry based on laboratory Investigations and quality control:

Critical / alert values

American Diabetes Association (ADA) Standards of Medical Care in Diabetes

(yearly update); HBA1C testing: NGSP

Case studies to review

Quality control for clinical chemistry in laboratory

#### **Recommended Texts**

- 1. ThomasM.Devlin (2014) Textbook of Biochemistry with Clinical Correlations (7th ed). John Wiley & Sons.
- 2. Montgomery R, Conway TW, Spector AA (1996), Biochemistry: A Case-Oriented Approach (6th ed), Mosby Publishers, USA.
- 3. Tietz Fundamentals of Clinical Chemistry and Molecular Diagnostics (2018) (8th ed), Saunders.
- 4. Dinesh Puri, (2020) Text book of Biochemistry: A clinically oriented approach 4th Edition, Elsevier.
- 5. M.N.Chatterjee and Rana Shinde (2012). Textbook of Medical Biochemistry (8th ed), Jaypee Brothers Medical Publishers.
- 6. Clinical Case Discussion In Biochemistry A Book On Early Clinical Exposure (ECE), Poonam Agrawal, 2021, CBS Publishers & distributors pvt. Ltd.

#### **Method of Evaluation:**

Test I	Test II	Assignment	Seminar	End Semester Examination	Total
5	10	5	5	75	100

#### **Methods of Assessment:**

**Recall (K1) - Simple definitions, MCQ, Recall steps, Concept definitions.** 

**Understand/Comprehend (K2) -** MCQ, True/False, Short essays, Concept explanations, Short summary or overview.

**Application (K3) -** Suggest idea/concept with examples, Observe, Explain.

**Analyse** (**K4**) - Finish procedure in stepwise manner, Differentiation between various ideas, Map knowledge.

**Evaluate (K5)** - Longer essay/ Evaluation essay, Critique or justify with pros and cons.

Create (K6) - Check knowledge in specific or offbeat situations, Discussion, Debating, Presentation.

#### **Mapping with Programme Outcomes:**

	PO 1	PO 2	PO 3	PO 4	PO 5	PO 6	PO 7	PO 8	PO 9	PO 10
CO 1	S	S	M	S	S	S	S	M	M	S
CO 2	S	M	S	M	S	S	S	M	M	M
CO 3	$\mathbf{S}$	S	S	S	S	$\mathbf{M}$	S	S	M	M
CO 4	S	M	M	M	S	M	S	S	S	M
CO 5	S	M	S	M	S	S	S	S	S	S

S-Strong M-Medium

L-Low

Cour	se / Course Code:	se / Course Code: CORE COURSE – VI / 23P3B11							
Title	e of the Course: GENE EDITING, CELL AND GENE THERAPY								
Sen	nester / Credits:	III/5							
Pre-r	requisites, if any:	To introduce students molecular basis of cell gene therapy;							
		viral and nonviral gene transfer techniques and gene therapy							
		applications in hereditary and acquired diseases.							
Cou	ırse Objectives:	1. To train the student in techniques related to the molecular							
		basis of genetic diseases and to incorporate skills essential							
		for various types ofsequencing.							
		2. To inculcate practical knowledge on comparing the animal							
		models used to model genetic diseases.							
		3. To introduce and also elaborate knowledge about wide							
		varieties of vectors and their features in addition to their							
		applications and to identify the viral and nonviral gene							
		transfer techniques.							
		4. To educate about the characteristics of cell culture,							
		therapeutic strategies in gene therapy with relevant							
<u> </u>	safety/ethics involved and patents aswell.								
Col	urse Outcomes:	After completion of the course, the students should be able to:							
		CO1: Ability to read, and evaluate scientific articles within the							
		subjects of immune therapy, gene therapy and cell							
		therapy (K1 & K2).							
		CO2: Toclone gene of their interest for several downstream							
		purposes							
		witharobustcomprehensionaboutwidevarietyofapplicableg							
		ene delivery vectors (K1, K2 &K5).							
		CO3: Be able to provide examples of diseases that can be treated with immune therapy, gene therapy and cell							
		therapy (K2, K3 & K4).							
		CO4: To identify knowledge gaps and need for further research							
		within their chosen topic of immune therapy, gene							
		therapy or cell therapy (K2, K4 & K5).							
		CO5: To critically discuss and reflect on ethical and social							
		aspects of using immune, gene or cell therapy. The							
		student will be persuaded to contemplate on upcoming							
		technologies for futuristic benefits (K2, K5 & K6).							
		Units							
	=	of gene editing, DNA repair mechanisms, Double strand DNA							
	breaks, Nonhomolo								
	_	eases for gene editing, Meganucleases, Zinc-Finger nucleases,							
	-	tor-Like Effector Nucleases (TALEN), CRISPR-Cas systems,							
	gene editing using (	CRISPR-Cas, drawbacks and major challenges to present gene							

	editing techniques, gene editing for human disease therapy.						
II							
	Gene and cell therapy: Basics of Gene and cell therapy, types of gene therapy, gene therapy strategies, therapeutic targets for gene therapy, choice of the therapeutic target, administration routes, delivery systems, expression of transgene, persistence of the gene therapy, cell targeting, immunological response to the therapy, ethical and legal issues, concerns about gene and cell therapy.  Vectors for Gene therapy: Non-viral and viral vectors for gene therapy, Physical methods of gene delivery, Polymer, Lipid and inorganic material based chemical systems for gene delivery, Viral vectors, Lentiviral, Adenoviral, Adeno-associated virus, Herpes Simplex virus, vaccinia, baculoviral vectors for gene delivery, choice of viral vector and oncolytic virus. Gene therapy applications, Gene therapy for cancer, suicide and oncolytic gene therapy.						
III							
	suicide and oncolytic gene therapy.						
IV							
V							
•							
	Genetically Modified Stem Cells in Experimental Gene Therapies. Technological						
	Reading List (Print and Online)						
	1. Stem Cell Biology, Daniel Marshak, Richard L. Gardener and David Gottlieb, Cold						
	Spring Harbour Laboratory Press.						
	2. Stem cell biology and gene therapy, Booth C., Cell Biology International,						
	Academic Press.						
	3. Stem Cell and Gene-Based Therapy: Frontiers in Regenerative Medicine,						
	Alexander Battler.						
	Self-Study						
	1. Applications of gene editing strategies						
<u> </u>	2. CART therapy for Cancer						
	Recommended Texts  1. An Introduction to Human Molecular Genetics (2nd Edition), J.J. Pasternak, 2005.						
	2. An Introduction to Molecular Medicine and Gene Therapy 1st Edition by Thomas F.						
	KresinaUpadhyay, S. K. (Eds.). (2021).						
	3. Human Molecular Genetics (4th Edition), Tom Strachan & Andrew Read, 2010.						
	4. Stem Cells Handbook: Stewart Sell, Humana Press; Totowa NJ, USA; Oct. 2003.						

Test I	Test II	Assignment	Seminar	End Semester Examination	Total
5	10	5	5	75	100

## **Methods of Assessment:**

Recall (K1) - Simple definitions, MCQ, Recall steps, Concept definitions.

**Understand/ Comprehend (K2) -** MCQ, True/False, Short essays, Concept explanations, Short summary or overview.

**Application (K3) -** Suggest idea/concept with examples, Observe, Explain.

**Analyse** (**K4**)- Finish procedure in stepwise manner, Differentiation between various ideas, Map knowledge

Evaluate (K5) - Longer essay/ Evaluation essay, Critique or justify with pros and cons.

**Create (K6)**- Check knowledge in specific or offbeat situations, Discussion, Debating, Presentation

# **Mapping with Programme Outcomes:**

	PO 1	PO 2	PO 3	PO 4	PO 5	PO 6	PO 7	PO 8	PO 9	PO 10
CO 1	S	L	M	S	M	M	M	M	M	M
CO 2	S	S	S	S	M	M	M	M	M	S
CO 3	S	M	S	S	M	S	S	S	S	S
CO 4	S	L	M	M	M	M	S	M	M	S
CO 5	S	S	S	S	S	S	S	S	S	S

Cour	rse / Course Code:	CORE COURSE –VII / INDUSTRIAL MODULE /23P3B12
Tit	le of the Course:	INDUSTRIAL MICROBIOLOGY
	nester / Credits:	III/5
Pre-	requisites, if any:	Basic Knowledge of Microbiology and microbial techniques.
Co	urse Objectives:	1. To gain knowledge of the structure, classification and use of
		microorganisms in various industries.
		2. To know various fermenter designs, culture systems and the
		<ul><li>application of fermentation process in industry.</li><li>3. To understand the production and purification of fermented</li></ul>
		products and their industrial applications.
		4. Understand the basic concepts of food and agricultural
		microbiology.
Co	ourse Outcomes:	CO1: Students will be able to understand the structure and
		classification of microorganisms (K2, K4).
		CO2: Gain knowledge of the uses of microorganisms in various
		industrial applications (K3, K4).
		CO3: Understand the concepts of fermentation process, harvest and recovery (K1, K5).
		CO4: Students will know the types of microbial fermentation
		processes and their applications in pharmaceutical industry
		(K2, K3).
		CO5: Students will learn about the use of microorganisms in
		beverages, diary and food industries (K3, K6).
		Units
I		eteria, fungi and viruses and their classification. Types and
		microorganisms used in Industry (a) Food Industry (b) Chemical
	Industry (c) Pharm	
II		l principles of microbial fermentation techniques – application in
	•	maceutical Biochemistry. Fermentation – types, techniques, design
	-	ermenters including addition of medium. Types and characteristics of
	•	environmental conditions required for the growth and metabolism of
	•	pharmaceutically important microbes. Sterilization methods in
		niques, air, gas, culture medium sterilization. Steam-filtration and
	fermentations, Ant	and constituents of fermentative culture medium and conditions of
III	· ·	imation of products of fermentation- Production of ethanol, acetic
111	•	tone, butanol and citric acid by fermentation. Production of Enzymes-
		, lipase, Production of pharmaceuticals by fermentation—penicillin,
		acycline, riboflavin, vitamin B12.Beverages-wine, beer and malt
	beverages.	acycline, moonavin, vitainin biz.beverages-wille, beer and matt
IV		gy: Production of dairy products-bread, cheese and yoghurt
		heir types). Food bore diseases- Bacterial and Non- Bacterial. Food
		nciples–Physical methods: temperature (low, high, canning, drying),
	preservation 1111	nerpies Injoien medicus. temperature (10 w, 111gh, emining, trying),

irradiation, hydrostatic pressure, high voltage pulse, microwave processing and aseptic
packaging, Chemical methods - salt, sugar, organic acids, SO <sub>2</sub> , nitrite and nitrates,
ethylene oxide, antibiotics and bacteriocins.
Agricultural Microbiology: General Properties of soil, microorganisms in soil -
decomposition of organic matter in soil. Biogeochemical cycles, nitrogen fixation,
Production of bio fertilizers and its field applications – Rhizobium, azotobacter, blue
green algae, mycorrhizae, azospirilium, Production of biofuels (biogas- methane), soil
inoculants.
Self-Study
Micro-organisms in food processing and pharma industries
Upstream and Downstream processes in Biopharma
Reading List (Print and Online)
1. Industrial biotechnology:https://nptel.ac.in/courses/102/105/102105058/
2. Bioreactors:https://nptel.ac.in/courses/102/106/102106053/
3. Food Microbiology:https://nptel.ac.in/courses/126/103/126103017/
4. Agriculture Microbiology:https://www.youtube.com/watch?v=f7UXyVImZ_c
Recommended Texts
1. Food Microbiology: An Introduction: 4 <sup>th</sup> edition, Matthews KR, Kniel KE,
Montville TJ; American Society for Microbiology.
2. Food, Fermentation and Micro-Organisms, 2 <sup>nd</sup> edition, Charles, BW;Blackwell
Science Ltd.
3. Microbiology. 5th edition, Pelczar MJ, Chan ECS and Krieg NR; McGraw Hill
BookCompany. 4. Text book of Microbiology: 11 <sup>th</sup> edition, Ananthanarayanan R and Paniker
4. Text book of Microbiology: 11 edition, Ananthanarayanan R and Paniker CKJ;Universities Press (India) Pvt.Ltd.
5. Food Microbiology, 3 <sup>rd</sup> edition,FrazierWC and Westhoff DC; Tata McGraw Hill
Publishing Company Ltd, NewDelhi.

Test I	Test II	Assignment	Seminar	End Semester Examination	Total
5	10	5	5	75	100

6. New Methods of Food Preservation: 1st edition, Gould GW; Springer Manual of

Industrial Microbiology and Biotechnology: 3rd edition, Baltz.

## **Methods of Assessment:**

Recall (K1) - Simple definitions, MCQ, Recall steps, Concept definitions.

**Understand/ Comprehend (K2) -** MCQ, True/False, Short essays, Concept explanations, Short summary or overview.

**Application (K3) -** Suggest idea/concept with examples, Solve problems, Observe, Explain.

**Analyse (K4) -** Problem-solving questions, Finish a procedure in many steps, Differentiate between various ideas.

**Evaluate (K5) -** Longer essay/ Evaluation essay, Critique or justify with pros and cons.

Create (K6) - Check knowledge in specific or offbeat situations, Discussion.

# **Mapping with Programme Outcomes:**

	PO 1	PO 2	PO 3	PO 4	PO 5	PO 6	PO 7	PO 8	PO 9	PO 10
CO 1	S	S	M	S	S	S	M	M	S	S
CO 2	S	M	S	S	M	S	S	M	M	M
CO 3	S	M	L	S	M	M	S	S	M	S
CO 4	M	S	S	S	L	M	S	M	S	M
CO 5	S	S	M	S	S	M	M	S	S	S

Course / Course Code:	CORE PRACTICAL – III / 23P3BP3
Title of the Course:	LABORATORY COURSE ON CLINICAL BIOCHEMISTRY
Semester / Credits:	III / 4
Pre-requisites, if any:	Knowledge on basic principles, Instrumentation of Biochemical techniques and metabolic reactions.
Course Objectives:	<ol> <li>To instill skill in students enabling them to apprehend the wider knowledge about principles and techniques to be employed for the investigation of biological samples, clinical approach, normal values of biochemical constituents and clinical interpretations.</li> <li>To inculcate the knowledge of collection, preservation of blood sample and learning various hematological parameters and their significance.</li> <li>To perform experiments to assess liver functions. And also to study the marker enzymes of liver</li> <li>To evaluate lipid profile and assess their relation to cardiac function.</li> <li>To perform experiments to estimate blood glucose and glycosylated hemoglobin.</li> <li>To perform urine analysis, estimate BUN and clearance test to assess renal function.</li> <li>To learn basic immune technniques antigen—antibody reactions.</li> <li>To perform data analysis in using MS Excel</li> <li>To introduce visit to hospital so that students may be aware of Phleobotomy, Collection and storage of specimen, Good laboratory practices, Automation and current methods adopted in the diagnostic labs</li> </ol>
Course Outcomes:	After completion of the course, the students should be able to: CO1: The student will be able to acquire knowledge and skill in hematology techniques. They will get familiar with methods and knowledge to interpret the electrolyte concentration in serum (K1 to K5).  CO2: The student will be able to assess the Liver Function and interpret the biochemical investigation in a given clinical situation (K1 to K5).  CO3: Skill to perform the Renal function test to assess the function of Kidney and report the abnormal parameters with reference range will be achieved by the student (K1 to K5).  CO4: To estimate the blood glucose content and lipid profile, to evaluate the alterations and record the observation in accordance to reference range will be acquired by the

	student (K1 to K6).						
	CO5: The Group Experiments will support them to acquire						
	practical skills to work in health care sector and assist then						
	to understand the automation process in clinical labs						
	(K1 to K6).						
	Units						
I	Haematology:						
	RBC count, WBC count – total and differential count, ESR, PCV, MCV. Bleedin time, Clotting Time, Grouping of blood and Rh typing and Estimation of hemoglobin.						
II	Liver function test:						
	Estimation of bilirubin – direct and indirect. Estimation of plasma protein, A/G ratio.						
	Thymol turbidity test, ProthrombinTime (PT), Assay of serum glutamate oxaloaceta transaminase, alkaline phosphatase, Gamma-glutamyltransferase (GGT).						
III	Renal function test:						
***	Collection and Preservation of Urine sample						
	Qualitative tests for normal and pathological components of urine.						
	BUN: Estimation of blood Urea, creatinine, and uric acid.						
IV	Estimation of blood glucose by orthotoluidine, Estimation of calcium and iron.						
	Estimation of cholesterol by Zak's method, estimation of triglycerides.						
V	Group Experiments						
	Antigen – Antibody Reaction - HCG kit method, RA kit method.						
	Phlebotomy – Venipuncture, Different techniques of venipuncture.						
	Collection of blood, Serum or Plasma separation and Storage.						
	Automation in Clinical Biochemistry – Autoanalyser and Semiautoanalyser.  Reading List (Print and Online)						
	1. https://www.researchgate.net/publication/260182512.						
	Practical Manual in Biochemistry and Clinical Biochemistry.						
	2. https://main.icmr.nic.in/sites/default/files/upload_documents/GCLP Guidelines 2020						
	•						
	Final.pdfhttps://www.westgard.com/clia.html						
	3. https://www.researchgate.net/publication/263929434, Biochemistry						
	4. https://ucms.ac.in/Lectures-C-2020/Renal%20function%20Tests%20-%20PPT.pdf						
	5. https://youtu.be/i2PfjEks4GQ						
	6. https://www.euro.who.int/data/assets/pdf file/0005/268790/WHO-guidelines-on-						
	drawing-blood-best-practices-in-phlebotomy-Eng.pdf						
	Self-Study						
	1. Laboratory handling of human biological specimen						
	2. Automation in Clinical Biochemistry						
	Recommended Texts						
	1. Practical Clinical Biochemistry- Varley's by Alan H Gowenlock, published by CBS						
	Publishers and distributors, India Sixth Edition, 1988.						
	2. Manipal Manual of Clinical Biochemistry (For Med.Lab.and Msc Stud.) 2013 (4)						
	Edition).						
	3. Case Oriented Approach in Biochemistry-Dr. Rajesh Kawaduji Jambhulkar, Dr.						
	Abhijit D. Ninghot: 2019 First Edition.						

- 4. Medical Lab Technology Vol I& II, Kanai L Mukerjee, New Delhi: Tata Mcgraw Hill Publishing Company, 1996.
- 5. Practical Biochemistry Plummer, New Delhi: Tata Mcgraw Hill Publishing Company, 2000.
- 6. Introductory Practical Biochemistry S.K. Sawhney, Randhir Singh, 2nd ed, 2005.

Test I	Test II	End Semester Examination	Total	Grade
20	20	60	100	

#### **Methods of Assessment:**

Recall (K1) - Simple definitions, MCQ, Recall steps, Concept definitions.

**Understand/ Comprehend (K2)** - MCQ, True/False, Short essays, Concept explanations, Short summary or overview.

**Application (K3)** - Suggest idea/concept with examples, Solve problems, Observe, Explain.

**Analyse** (**K4**) – Problem-saving questions, Finish a procedure in many steps, Differentiate between various ideas.

Evaluate (K5) - Longer essay/ Evaluation essay, Critique or justify with pros and cons

Create (K6) – Check knowledge in specific or offbeat situations. Discussion.

## **Mapping with Programme Outcomes:**

	PO 1	PO 2	PO 3	PO 4	PO 5	PO 6	PO 7	PO 8	PO 9	PO 10
CO 1	S	S	S	S	M	S	L	S	M	S
CO 2	S	S	S	S	M	S	L	S	M	S
CO 3	S	S	S	S	M	S	M	S	M	S
CO 4	S	S	S	S	M	S	M	S	S	S
CO 5	S	S	S	S	S	S	S	S	S	S

Cou	rse / Course Code:	ELECTIVE COURSE – V / 23P3B13EC					
Ti	tle of the Course:	BIOSTATISTICSAND DATA SCIENCE					
Se	mester / Credits:	III / 4					
Pre-	requisites, if any:	Basic knowledge of Statistics and Computer Applications.					
Co	ourse Objectives:	1. To summarize the data and to obtain its salient features from					
		thevastmass of original data.					
		2. To understand the concept of various measures of					
		dispersion.					
		3. To understand the concepts of sampling and learning test					
		of significance.					
		4. To understand the concept of various attributes and relate to					
		biological studies.					
		5. To gain knowledge in SPSS, a software package which					
		gives aperfect graphical representation and appropriate					
		result for thedata that has been entered.					
Co	ourse Outcomes:	After completion of the course, the students should be able to:					
		CO1: Concepts of statistical population and sample, variables					
		and attributes. Tabular and graphical representation of data based on variables (K1, K2& K3).					
		CO2: Conditions for the consistency' and criteria for the					
		independence of data based on attributes. Measures					
		central tendency, Dispersion, Skewness and Kurtosis					
		(K1, K2& K3).					
		CO3: Learning different sampling methods and analysing					
		statistical significance (K1, K2, K3& K4) CO4: Understanding students t test, ANOVA, Chi square test					
		to analyse the significance of various research (K1, K2,					
		K3& K4).					
		CO5: Learning on data science, algorithm for machine					
		learning, artificial intelligence and big data, their					
		applications in clinical and pharma domain (K1, K2,					
		K3, K4& K6).					
	T	Units					
Ι		l and clinical experiments – Collection of data in experiment-					
	Primary and secondary data. Methods of data collection. Classification and						
	tabulation. Different forms of diagrams and graphs related to biological studies						
	Measures of Averages- Mean, Median, and mode. Use of these measures in						
	biological studies.						
II	_	ersion for biological characters— Quartile deviation, Mean					
	*	deviation and coefficient of variation. Measures of skewness and					
		on and regression – Rank correlation – Regression equation. sed on biochemical data.					
	Simple problems bas	ou on oroenement unin.					

III	Basic concepts of sampling- Simple random sample stratified sample and systemic sampling. Sampling distribution and standard error. Test of significance based on large samples. Test for mean, difference of means, proportions and equality of proportions.						
IV	Small sample tests – Students't' test for mean, difference of two way means, tests for correlation and regression coefficients. Chi-square test for goodness of a non independence of attributes. F test for equality of variances. ANOVA- one way and two way. Basic concept related to biological studies						
V	Introduction to Data Science, Definition of data science, importance, and basic applications, Machine Learning Algorithms, Deep Learning, Artificial Neural Networks and their Application, Reinforcement Learning, Natural Language Processing Artificial Intelligence (AI), Data Visualization, Data Analysis, Optimization Techniques, Big Data, Predictive Analysis. Application of AI in medical, health and pharma industries.						
	Reading List (Print and Online)  1. https://www.ibm.com/docs/en/SSLVMB_28.0.0/pdf/Accessibility.pdf  2. https://pure.tue.nl/ws/portalfiles/portal/19478370/20160419_CO_Mzolo.pdf  3. https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5453888/  4. https://home.ubalt.edu/ntsbarsh/excel/excel.htm  5. https://students.shu.ac.uk/lits/it/documents/pdf/analysing_data_using_spss.pdf  6. https://www.ibm.com/support/pages/ibm-spss-statistics-28-documentation						
	Self-Study						
	<ol> <li>Simple problems on probability, theoretical distributions, hypothesis testing</li> <li>Relationship between mean, median and mode pros and cons of the measures of central tendency and deviation</li> </ol>						
	Recommended Texts						
	1. Zar, J.H. (1984) "Bio Statistical Methods", Prentice Hall, International Edition						
	2. Sundar Rao P. S.S., Jesudian G. & Richard J. (1987), "An Introduction to						
	Biostatistics", 2nd edition, Prestographik, Vellore, India.						
	3. Warren, J; Gregory, E; Grant, R (2004), "Statistical Methods in Bioinformatics", 1st						
	edition,Springer						
	4. Milton, J.S. (1992),. "Statistical methods in the Biological and Health Sciences", 2nd						
	edition,Mc Graw Hill,						
	5. Rosner,B (2005), "Fundamentals of Biostatistics", Duxbury Press						
	6. Introducing Data Science, Davy Cielen, Anro DB Meysman, Mohamed Ali.						

Test I	Test II	Assignment	Seminar	End Semester Examination	Total
5	10	5	5	75	100

# **Methods of Assessment:**

Recall (K1) - Simple definitions, MCQ, Recall steps, Concept definitions.

**Understand/ Comprehend (K2) -** MCQ, True/False, Short essays, Concept explanations, Short summary or overview.

Application (K3) - Suggest idea/concept with examples, Solve problems, Observe, Explain

**Analyse (K4) -** Problem-solving questions, Finish a procedure in many steps, Differentiate between various ideas

**Evaluate (K5)** - Longer essay/ Evaluation essay, Critique or justify with pros and cons **Create (K6)** - Check knowledge in specific or off beat situations, Discussion, Presentations

# **Mapping with Programme Outcomes:**

	PO 1	PO 2	PO 3	PO 4	PO 5	PO 6	PO 7	PO 8	PO 9	PO 10
CO 1	S	S	S	S	M	S	L	S	S	S
CO 2	S	S	S	S	M	S	L	S	S	S
CO 3	S	S	S	S	S	S	M	S	S	S
CO 4	S	S	S	S	S	S	M	S	S	S
CO 5	S	S	S	S	S	S	M	S	S	S

Course / Course Coue:	rse / Course Code: SKILL ENHANCEMENT COURSE [SEC] – II/ 23P3B14SEC								
Title of the Course:	MOLECULAR BASIS OF DISEASES AND THERAPEUTIC								
	STRATEGIES ts: III / 2								
Semester / Credits:	III / 2								
Pre-requisites, if any:	Knowledge of Human Physiology, Metabolism and Clinical								
	Biochemistry.								
Course Objectives:	1. To understand the concepts of the mechanisms involved in regulation of blood sugar and management of diabetes								
	mellitus								
	2. To gain in-depth knowledge of the mechanisms of cancer and of tumor metastasis								
	3. The student will review the basic organization of the central								
	and peripheral nervous system that coordinate the sensory and								
	motor functions of the body. In addition, the student will								
	explore impaired features underlying the major								
	neuropathological complications.								
	<ul><li>4. To gain knowledge in renal diseases</li><li>5. To understand the mechanisms involved in cardiac disorders</li></ul>								
Course Outcomes:									
Course outcomes.	understand								
	_								
	interventions.								
	CO5: A thorough knowledge on the experimental models of non-								
	communicable diseases that will be applied for future								
	research or project dissertation. An in-depth knowledge on								
	development of drugs against non-communicable diseases.								
	Units								
I Mechanism of blo	ood sugar regulation in human body. Pathophysiology of Type I and II								
	s – investigation methods for the diagnosis of diabetes. Nutritional								
	Complications related to diabetes – Diabetic cardiovascular disease, retinopathy,								
-	thy and nephropathy. Cellular and molecular mechanism of development of								
	ement of Type I and Type II diabetes, drugs for the treatment of								
diabetes.									
II Biology of cand	cer: Overview of hallmarks of cancer. Tumorigenesis, Tumor								
	mechanism of Metastasis. Proto-oncogene to oncogene. Oncogene-								
1 2	ly. Tumor suppressor gene-Rb and p53 pathway in cancer. Diagnosis-								
diabetes, Diabete care. Complication neuropathy and rediabetes- Manage diabetes.  II Biology of canon progression and rediabetes.	CO1: Overall view about the complications of diabetes mellitus and its management.  CO2: Comprehensive understanding of the concepts of cancer biology and implicating the theoretical concepts for further research.  CO3: Understand and appreciate the pathophysiology of conditions affecting the nervous system.  CO4: A thorough knowledge of renal and cardiac diseases with emphasis related to mechanistic aspects and therapeutic interventions.  CO5: A thorough knowledge on the experimental models of noncommunicable diseases that will be applied for future research or project dissertation. An in-depth knowledge on development of drugs against non-communicable diseases.  Units  Ood sugar regulation in human body. Pathophysiology of Type I and II is – investigation methods for the diagnosis of diabetes. Nutritional instrelated to diabetes – Diabetic cardiovascular disease, retinopathy, nephropathy. Cellular and molecular mechanism of development of the treatment of the core: Overview of hallmarks of cancer. Tumorigenesis, Tumor mechanism of Metastasis. Proto-oncogene to oncogene. Oncogene-								

	Non-invasive imaging techniques, Tumor diagnosis, Interventional radiology, New							
	imaging technique, Molecular techniques in cancer diagnosis treatment of cancer-							
	surgery, radiotherapy, chemotherapy, hormonal treatment, and biological therapy.							
	Introduction to personalized medicine.							
TIT								
III	Brain- neuronal network- memory- Neurogenerative diseases- Parkinson and							
	Alzheimer Disease- molecular understanding of the neurodegenerative diseases-							
	treatment modalities.							
IV	Acute and chronic renal failure, glomerular diseases–glomerulonephritis, nephritic							
	syndrome, diabetes insipidus, diagnosis of kidney disease.							
V	Introduction to cardiovascular diseases, Lipids and lipoproteins in coronary heart							
	disease-cardiac enzymes, Molecular changes during cardiac remodeling – hypertrophy							
	of hearts – heart failure- treatment modalities.							
	Reading List (Print and Online)							
	1. The Biochemical basis of disease:2018,Barr AJ; Portland Press							
	2. Biochemical Basis of Diseases							
	3. https://www.biologydiscussion.com/diseases-2/biochemical-basis-of-							
	diseases/44276							
	Recommended Texts							
	1. Wills' Biochemical Basis of Medicine: 2 <sup>nd</sup> edition, Thomas H, Gillham B; Elsevier							
	2. Molecular Biochemistry of Human Diseases, 2021, Feuer G, de la Iglesia F; CRC							
	Press.							

Test I	Test II	Test II Assignment		Test II Assignment Seminar		End Semester Examination	Total
5	10	5	5	75	100		

## **Methods of Assessment:**

Recall (K1) - Simple definitions, MCQ, Recall steps, Concept definitions.

**Understand/ Comprehend (K2) -** MCQ, True/False, Short essays, Concept explanations, Short summary or overview.

**Application (K3) -** Suggest idea/concept with examples, Suggest formulae, Solve problems, Observe, Explain.

**Analyse (K4) -** Problem-solving questions, Finish a procedure in many steps, Differentiate between various ideas, Map knowledge.

Evaluate (K5) - Longer essay/ Evaluation essay, Critique or justify with pros and cons.

# **Mapping with Programme Outcomes:**

	PO 1	PO 2	PO 3	PO 4	PO 5	PO 6	PO 7	PO 8	PO 9	PO 10
CO 1	S	S	S	M	M	S	S	S	S	S
CO 2	S	M	S	L	M	M	M	M	M	S
CO 3	S	S	M	L	S	S	M	M	S	M
CO 4	S	M	M	M	M	M	S	S	M	S
CO 5	S	S	M	M	S	M	M	M	S	S

Course / Course Code:		CORE COURSE – VIII / 23P4B15						
Titl	e of the Course:	PHARMACEUTICAL BIOCHEMISTRY						
Ser	nester / Credits:	IV / 5						
Pre-	requisites, if any:	The student should have a basic knowledge of drug discovery and development. Student should possess basic knowledge bioinformatics to understand and correlate the drug development process.						
Cour	se Objectives:	<ol> <li>To understand the different types of bioinformatic tools for drug discovery.</li> <li>To get an overview of how different bioinformatic toolsaid in the process of target identification, drug screening and quantitative structure activity relationship.</li> <li>To assimilate the involvement of different metabolic pathways involved in drug metabolism and correlate their involvement in elimination process</li> </ol>						
		<ul><li>4. To understand the biochemical basis of drug action at the target tissue.</li><li>5. To understand different phases in drug clinical trials and its assessment.</li></ul>						
Co	ourse Outcomes:	<ul> <li>After completion of the course, the students should be able to:</li> <li>CO1: To understand and explain the basic concepts of drug discovery and drug development process.</li> <li>CO2: To review the different software and computational tools which aid in the design of drugs and its rationalization.</li> <li>CO3: To analyze the different stages of the drug discovery process with the target &amp; hit identification, assays for drugscreening and preclinical studies.</li> <li>CO4: To understand the various phases of the clinical trialsand the method of conduct of clinical trials.</li> </ul>						
		Units						
I	Drug discovery and development, drug target identification and validation, H identification, General principles of screening, correlations between various animal models and human situations, Correlation between in-vitro and in-vivo screens; Special emphasis on cell-based assay, biochemical assay, radiological binding assay Pharmacological assay, In vitro, In vivo & Ex-vivo experiments, lead optimization preclinical studies.							
II	Bioinformatics approaches for drug development:  Identification of potential molecules, chemical compound library preparation, Identification of target in pathogen, Ligand & protein preparation, Molecular docking, Binding free energy estimation, High throughput virtual screening, Docking protocol validation and enrichment analysis, Single point energy calculation, Pharmacokinetics and Pharmacodynamics, ADME & toxicity prediction, Molecular dynamic simulation,							

Rule of three and five, Lipinsky rule, Pharmacophore development, Quantitative structure activity relationship, 3D-QSAR, Techniques of developing a pharmacophore map covering both ligand based and receptor based approaches.

# **III** Drug metabolism & interactions:

Drug-receptor interactions, receptor theories and drug action, Xenobiotics, xenobiotics phases (Phase-I, Phase-II and Phase-III), role of cytochrome P450 oxidases and glutathione S-transferases in drug metabolism, factors affecting drug metabolism, Enzymes as a drug target, Kinase inhibitors, ATPase inhibitors, drug protein interaction, DrugDNA interaction. Basic ligand concepts-agonist, antagonist, partial agonist, inverse agonist, efficiency and potency. Forces involved in drug-receptor complexes. Receptor classification – the four super families. Receptor binding assays- measurement of Kd, Bmax and  $IC_{50}$ .

- IV Biochemical mode of action of antibiotics- penicillin and chloramphenicol, actions of alkaloids, antiviral and antimalarial substances. Biochemical mechanism of drug resistance- sulphonamides. Drug potency and drug efficacy. General principles of chemotherapy: chemotherapy of parasitic infections, fungal infections, viral diseases. Introduction to immunomodulators and chemotherapy of cancer.
- V Clinical trials (Phase-I, Phase-II, Phase-III and Phase-IV clinical trial). Main features of clinical trials, including methodological and organizational considerations and the principles of trial conduct and reporting. Key designs surrounding design, sample size, delivery and assessment of clinical trials.

## **Self-Study**

- 1. Examples of pharmaceutical development of a drug
- 2. Basic pharmacology of drug action and kinetics

# **Reading List (Print and Online)**

- 1. Textbook of Drug Design. Krogsgaard-Larsen, Liljefors and Madsen (Editors), Taylor and Francis, London UK, 2002.
- 2. Drug Discovery Handbook S.C. Gad (Editor) Wiley-Interscience Hoboken USA, 2005.

## **Recommended Texts**

- 1. Practical Application of Computer-Aided Drug Design, Ed. Charifson P., Marcel Dekker Inc.
- 2. 3D QSAR in Drug Design: Theory, Methods and Applications, Ed. Kubinyi H., Ledien.
- 3. Pharmaceutical Profiling in Drug Discovery for Lead Selection, Borchardt RT, Kerns, EH, Lipinski CA, Thakker DR and Wang B, AAPS Press, 2004.
- 4. Drug Discovery and Development; Technology in Transition. HP Rang. Elsevier Ltd 1st edition 2006.
- 5. Pharmacology in Drug Discovery. T. P. Kenakin. Elsevier, 1st Edition 2012.

Test I	Test II	Assignment	Seminar	End Semester Examination	Total	
5	10	5	5	75	100	

#### **Methods of Assessment:**

**Recall** (K1) – Simple definitions, MCQ, Recall steps, Concept definitions.

**Understand/Comprehend (K2) -** MCQ, True/False, Short essays, Concept explanations, Short summary or overview.

**Application (K3) -** Suggest idea/concept with examples, Solve problems, Observe, Explain.

**Analyse** (**K4**) – Problem-saving questions, Finish a procedure in many steps, Differentiate between various ideas.

**Evaluate (K5)** - Longer essay/ Evaluation essay, Critique or justify with pros and cons.

Create (K6) – Check knowledge in specific or offbeat situations. Discussion.

# **Mapping with Programme Outcomes:**

	PO 1	PO 2	PO 3	PO 4	PO 5	PO 6	PO 7	PO 8	PO 9	PO 10
CO 1	S	M	S	M	S	M	M	S	S	S
CO 2	S	S	S	M	M	S	S	S	S	S
CO 3	S	S	S	L	S	M	M	S	S	M
CO 4	S	M	S	L	S	L	M	S	S	M
CO 5	S	S	S	L	S	M	M	S	S	S

Course / Course Code:	CORE PRACTICAL – IV/ 23P4BP4
Title of the Course:	LABORATORY COURSE ON NUTRITIONAL
	BIOCHEMISTRY AND BIOLOGY
Semester / Credits:	IV / 5
Pre-requisites, if any:	Knowledge on basic principles, instrumentation of biochemical
	techniques and applications.
Course Objectives:	1. To develop skill in students enabling them to apprehend the
Course Objectives.	wider knowledge about principles and techniques to be
	employed for the analysis of biological samples.
	2. To perform experiments to assess food quality and
	biochemical contents and create the awareness of food and its
	analysis.
	3. To understand the food analysis by both qualitative and
	quantitative methods and acquire knowledge on the
	sterilization techniques, media preparation and isolation of
	bacteria.
	4. To perform experiments to estimate pyruvate and tryptophan
	and learn principle and technique of agarose gel
	electrophoresis.
	5. To inculcate the knowledge of various isolation and
	purification techniques of biomolecules and achieve training
	in sub-cellular fractionation and to identify them by markers.
	6. To perform the isolation and identification of the organelles of a cell using differential centrifugation.
Course Outcomes:	On successful completion of this course, students should be able to:
Course Outcomes.	After completion of the course, the students should be able to:
	CO1: The student will be able to acquire knowledge and skill in
	isolation techniques. They will get familiar with methods and
	knowledge to interpret the biomolecules concentration in
	biological samples (K1 to K5).
	CO2: The student will be able to assess the quantity of biomolecules
	and interpret the biochemical investigation in a given
	biological samples (K1 to K5).
	CO3: Skill to perform the analysis of food adulterants in food
	materials and report the adulterants will be achieved by the
	student (K1 to K5).
	CO4: To study the microbial techniques - sterilization, Gram
	staining and media preparation and student will be acquired
	knowledge on good laboratory practices in microbiology
	laboratory (K1 to K6).
	CO5: The students will develop skill in analytical techniques like
	subcellular fractionation and Agarose gel electrophoresis and

	the group experiments will enable them to build learning							
	skills like team work, Problem solving, Communication							
	ability (K1, K2,K3,K4 & K6).							
	Units							
I	Isolation of Biomolecules							
	Isolation of casein from milk							
	Isolation of lactose from milk							
	Isolation of gluten from wheat flour							
	Isolation of starch from potato.							
	Isolation and estimation of glycogen from liver							
II	Quantitative Analysis							
	Estimation of reducing sugar in milk – Benedicts method							
	Estimation of iron content from biological sample							
	Estimation of albumin from the egg white.							
	Estimation of polyphenols							
	Determination of rancidity in edible oil							
	Estimation of pyruvate							
	Estimation of tryptophan.							
III	Analysis of Food Adulterants							
	Edible oil							
	Milk and milk products							
TX 7	Beverages, spices and condiments							
IV	Microbial Techniques  Sofety massyras and good laboratory practices in microbiology laboratory							
	Safety measures and good laboratory practices in microbiology laboratory							
	Sterilization, culture and inoculums preparation							
	Staining of bacteria – Gram Staining							
₹7	Preparation of media							
V	Group Experiments							
	Fractionation of sub-cellular organelles by differential centrifugation – Mitochondria							
	and nucleus							
	Identification of the separated sub-cellular fractions using marker enzymes (any one)							
	Agarose gel electrophoresis of genomic DNA							
	Reading List (Print and Online)							
	1. https://www.researchgate.net/publication/260182512.							
	Practical Manual in Biochemistry and Clinical Biochemistry.							
	2. https://main.icmr.nic.in/sites/default/files/upload_documents/GCLP Guidelines 2020							
	Final.pdfhttps://www.westgard.com/clia.html							
	3. https://www.researchgate.net/publication/263929434, Biochemistry							
	4. https://ijpsr.com/bft-article/determination-of-total-flavonoid-and-phenol-content-in-							
	mimusops-elengi-linn/?view=fulltext							
	5. https://skyfox.co/wp-content/uploads/2020/12/Practical-Manual-of-Biochemistry.pdf							
	Self Study							
	1. Moisture, ash and fiber contents of a food sample							
	2. Preparation of culture media and reagents							

#### **Recommended Texts**

- 1. Damodaran, S., Parkin, K.L. and Fennema, O. R. (2007). Fennema's Food Chemistry, fourth edition, published by CRC Press.
- 2. Meyer L.H. (2003). Food Chemistry, Reinhold Pub. Corp.
- 3. Nielsen, S.S. (2003). Food Analysis, Third Ed., Kluwer Academic/Plenum Publishers, New York.
- 4. Beedu Sashidhar Rao, Viay Deshpande. Experimental Biochemistry. I.K International Pvt Ltd.
- 5. Analytical techniques in Biochemistry and Molecular Biology; Katoch, Rajan. Springer (2011).
- 6. Basic Methods for the Biochemical Lab; Martin Holtzhauer, Springer, (2007).
- 7. Wilson and Walker's Principles and Techniques of Biochemistry and Molecular Biology, 8th Edn. Andreas Hoffman and Samuel Clockie, Ed., Cambridge University Press, (2018).
- 8. Biochemistry LabFax, Ed. J.A.A. Chambers and D. Rickwood, Blackwell Science, (1993).
- 9. Biochemical Techniques 87th Edn., John F. Roby & Bernard J White Waveland Press Inc. (1987).

#### **Method of Evaluation:**

Test I	Test II	End Semester Examination	Total	Grade
20	20	60	100	

#### **Methods of Assessment:**

**Recall (K1)** - Simple definitions, MCQ, Recall steps, Concept definitions.

**Understand/ Comprehend (K2)** - MCQ, True/False, Short essays, Concept explanations, Short summary or overview.

**Application** (K3) - Suggest idea/concept with examples, Solve problems, Observe, Explain.

**Analyse** (**K4**) – Problem-saving questions, Finish a procedure in many steps, Differentiate between various ideas.

Evaluate (K5) - Longer essay/ Evaluation essay, Critique or justify with pros and cons

Create (K6) – Check knowledge in specific or offbeat situations. Discussion.

# **Mapping with Programme Outcomes:**

	PO 1	PO 2	PO 3	PO 4	PO 5	PO 6	PO 7	PO 8	PO 9	PO 10
CO 1	S	S	S	S	M	S	L	S	M	S
CO 2	S	S	S	S	M	S	L	S	M	S
CO 3	S	S	S	S	M	S	M	S	M	S
CO 4	S	S	S	S	M	S	M	S	S	S
CO 5	S	S	S	S	S	S	S	S	S	S

Cours	se / Course Code:	ELECTIVE COURSE – VI / 23P4B16EC					
Title	e of the Course:	BIOCHEMICAL TOXICOLOGY					
Sem	nester / Credits:	IV / 4					
Pre-r	requisites, if any:	The student should have a basic knowledge of pharmacology of					
		drug action and understanding on their biochemical pathways.					
Cou	ırse Objectives:	1. To understand the detailed study of biochemical basis of					
		drugs and its toxicity, particularly their actions on living					
		systems.					
		2. To understand the relevance and methods to identify the					
		chemotherapeutic value of drug.					
		3. To understand the fundamentals of toxicology and dose-					
		response relationships.					
		4. To understand the toxicological drug testing procedures based					
		on in vitro and animal studies					
		5. To understandbiochemical pathways of drug toxicity and its					
	0.1	manifestation on vital organs.					
Cou	urse Outcomes:	On completion of this course, the student will be able					
		CO1: To appreciate and understand the role of toxicological					
		biomarkers to assess drug toxicities.					
		CO2: To conceive the role of disposition of drug in human system					
		and their metabolism and methodologies pertaining to toxicological studies.					
		CO3: To understand and evaluate the functions of different					
		organson drug disposition and associated drug toxicities.					
		CO4: To understand the toxicological response to foreign					
		compounds and their pharmacological, physiological and					
		biochemical effects.					
		CO5: To link the mechanism of toxicity and clinical symptoms					
		with underlying physiological disturbances.					
		Units					
I	Fundamentals of Toxicology and dose-Response Relationships:						
	Introduction Biomarkers Criteria of Toxicity New Technologies Evaluation of Toxicity						
		teractions; Dose Response; Measurement of Dose-Response; Relationships Linear					
	_	Hormesis; Hazard and Risk Assessment Duration and Frequency of					
	Exposure and Effe	ect.					
II		Toxic Responses: Disposition: Absorption, Sites of absorption,					
	distribution, Excre	etion; Metabolism: types of Metabolic change phase I reactions;					
	Phase 2 reactions;	control of Metabolism, Toxication vs. Detoxication.					

III	Toxicity testing; Test protocol, Genetic toxicity testing & Mutagenesis assay: In vitro							
	test systems: bacterial mutation tests-Reversion test, Ames test, Fluctuation test, and							
	Eukaryotic mutation test. In vivo test system Mammalian mutation test-Host mediated							
	assay and Dominant Lethal test. Biochemical basis of toxicity: Mechanism of toxicity:							
	Disturbance of excitable membrane function, Altered Calcium homeostasis, Covalent							
	binding to cellular macromolecules & genotoxicity, Tissue specific toxicity.							
IV	Toxic Responses to Foreign Compounds: Direct Toxic Action: Tissue Lesions;							
1 4								
	Mechanism and response in cellular toxicity, pharmacological, physiological and							
	Biochemical effects; Developmental Toxicology-Teratogenesis; Immunotoxicity							
	Genetic Toxicity; Chemical Carcinogenesis.							
V	Biochemical Mechanisms of Toxicity: Tissue Lesions: Liver Necrosis; kidney							
	Damage; Lung Damage, Liver damage, Cardiac damage; Neurotoxicity; Exaggerated							
	and Unwanted pharmacological effects; Physiological effects; Biochemical Effects:							
	Lethal Synthesis and Incorporation, Interaction with specific Protein Receptors;							
	Teratogenesis; Immunotoxicity; multi-Organ Toxicity.							
	Self-Study							
	Case studies to review							
	Reading List (Print and Online)							
	1. Preclinical Safety Evaluation of Biopharmaceuticals: A Science - Based Approach							
	to Facilitating Clinical Trials by Joy A. Cavagnaro							
	2. A Comprehensive Guide to Toxicology in Nonclinical Drug Development 2nd							
	Edition by Ali S. Faqi							
	Recommended Texts							
	1. Principles Of Toxicology by: Karen E Stine, Thomas M Brown 2006							
	Publisher. Crc Press.							
	2. Principles of Biochemical Toxicology by John A. Timbrell Publisher:							
	Informa Healthcare.							
	3. Environmental Toxicology by Sigmund F. Zakrzewski, (2002)							
	Publisher: Oxford University Press, USA.							

Test I	Test II	Assignment	Seminar	End Semester Examination	Total
5	10	5	5	75	100

# **Methods of Assessment:**

**Recall (K1)** – Simple definitions, MCQ, Recall steps, Concept definitions.

**Understand/Comprehend (K2) -** MCQ, True/False, Short essays, Concept explanations, Short summary or overview.

**Application** (**K3**) – Suggest idea/concept with examples, Solve problems, Observe, Explain.

**Analyse** (**K4**) – Problem-saving questions, Finish a procedure in many steps, Differentiate between various ideas.

**Evaluate (K5)** - Longer essay/Evaluation essay, Critique or justify with pros and cons.

Create (K6) – Check knowledge in specific or offbeat situations, Discussion

# **Mapping with Programme Outcomes:**

	PO 1	PO 2	PO 3	PO 4	PO 5	PO 6	PO 7	PO 8	PO 9	PO 10
CO 1	S	S	S	L	S	L	M	M	M	M
CO 2	M	M	S	M	M	L	M	S	S	S
CO 3	S	S	S	M	M	L	S	S	M	M
CO 4	S	M	S	M	M	M	S	S	M	M
CO 5	M	S	S	S	S	M	M	M	S	S

Course / Course Code:	SKILL ENHANCEMENT COURSE [SEC] – III/ 23P4B17SEC					
Title of the Course:	BIOSAFETY, LAB SAFETY AND IPR					
Semester / Credits:	IV / 2					
Pre-requisites, if any:	The student should have a basic knowledge of hazards associated					
	with the handling of biological agents and importance of					
	intellectual property from scientific research.					
Course Objectives:	1. To assimilate the hazards associated with the handling of					
	biological and chemical agents.					
	2. To understand how to protect from the hazards by the					
	implementation of various safety measures in biochemical					
	laboratories.					
	3. To implicate the importance of protecting the scientific					
	intellect by filing patent and understand the various offices for					
	filing and maintaining patents.					
	4. To understand the scope of patenting in biological research.					
	5. To create an awareness of ethics associated with used of					
	genetically modified organisms/cells and its rationale for use in					
	living organisms.					
Course Outcomes:	After completion of the course, the students should be able to:					
	CO1: To understand and implement various aspects of biosafety					
	and carry out risk assessment of products in biological					
	research CO2: Understand the basic concepts of ethics and safety that are					
	essential for different disciplines of science and procedures					
	involved and protection of intellectual property and related					
	rights.					
	CO3: To appreciate the intellectual property rights and its					
	implementation of on the invention related to biological					
	research.					
	CO4: To understand the statutory bodies that regulate the					
	property rights and its validity in various countries.					
	CO5: Critique the ethical concerns associated with modern					
	biotechnology processes and plan accordingly.					
I Biosafety: Historic	Units					
-	Biosafety: Historical background; introduction to biological safety cabinets; primary containment for biohazards; biosafety levels; recommended biosafety levels for					
	and infected animals; biosafety guidelines - government of India,					
	CGM, GEAC etc. for GMO applications in food and agriculture;					
	elease of GMOs; risk assessment; risk management and					
Communication, na	national regulations and international agreements.					

Laboratory safety - Chemical, electrical and fire hazards; handling and manipulating human or animal cells and tissues, toxic, corrosive or mutagenic solvents and reagents; mouth pipetting, and inhalation exposures to infectious aerosols, Safe handling of syringe needles or other contaminated sharps, spills and splashes onto skin and mucous membranes. Health aspects; toxicology, allergenicity, antibiotic resistance.

History of biosafety microbiology and molecular biology, Risk assessment, Personal protective equipment, Laboratory facilities and safety equipment, Disinfection, decontamination, and sterilization, Regulatory compliance, Laboratory security and emergency response and administrative controls.

- III Intellectual Property Rights (IPR): Introduction to patents, types of patents, process involved in patenting in India, trademarks, copyright, industrial design, trade secrets, traditional knowledge, geographical indications, history of national and international treaties and conventions on patents, WTO, GATT, WIPO, Budapest Treaty, Patent Cooperation Treaty (PCT) and TRIPS. Patent databases: Searching international databases; analysis and report formation. Indian Patent Act 1970; recent amendments; filing of a patent application; precautions before patentingdisclosure/non-disclosure; procedure for filing a PCT application. The patentability of microorganisms-claims, Characterization and repeatability disposition in the culture collections, legal protection for plants and other higher organisms, new plant varieties by rights, tissue culture protocols
- Patent filing and infringement: Patent application- forms and guidelines, fee structure, time frames; types of patent applications: provisional and complete specifications; PCT and convention patent applications, International patenting-requirement, financial assistance for patenting-introduction to existing schemes; Publication of patents-gazette of India, status in Europe and US. Research Patenting: Patenting by researchers and scientists-University/organizational rules in India and abroad. Detailed information on patenting biological products, Case studies on patents (basmati rice, turmeric, neem etc.), and patent infringement.

## V Bioethics:

Introduction to bioethics, human genome project and its ethical issues, genetic manipulations and their ethical issues, ethical issues in GMOs, foods and crops in developed and developing countries, environmental release of GMOs, ethical issues involved in stem cell research and use, use of animals in research experiments, animal cloning, human cloning and their ethical aspects, testing of drugs on human volunteers.

## **Self-Study**

- 1. Review of drug patent documents
- 2. Safety in biological research laboratories

# **Reading List (Print and Online)**

- 1. V. Shree Krishna, (2007). Bioethics and Biosafety in Biotechnology, New Age International Pvt. Ltd. Publishers. (Unit III, Unit IV and Unit V).
- 2. Deepa Goel, Shomini Parashar, (2013). IPR, Biosafety and Bioethics, Pearson. (Unit II)

- 3. R. Ian Freshney, 2016. Culture of Animal Cells: A Manual of Basic Technique and Specialized Applications, 6th Ed, John Wiley & Blackwell.
- 4. BAREACT, Indian Patent Act 1970 Acts & Rules, Universal Law Publishing Co. Pvt. Ltd., 2007. (Unit I)

## **Recommended Texts**

- Biosafety in Microbiological and Biomedical Laboratories, (2020), 6th Ed. (https://www.cdc.gov/labs/pdf/SF\_19\_308133-A\_BMBL6\_00-BOOK-WEB-final3.pdf)
- 2. Kankanala C., (2007), Genetic Patent Law & Strategy, 1st Edition, Manupatra Information Solution Pvt. Ltd.

#### **Method of Evaluation:**

Test I	Test II	Assignment	Seminar	End Semester Examination	Total	
5	10	5	5	75	100	

#### Methods of A

#### ssessment:

**Recall** (K1) – Simple definitions, MCQ, Recall steps, Concept definitions.

**Understand/Comprehend (K2) -** MCQ, True/False, Short essays, Concept explanations, Short summary or overview.

**Application**(**K3**) - Suggest idea/concept with examples, Solve problems, Observe, Explain.

**Analyse** (**K4**) – Problem-saving questions, Finish a procedure in many steps, Differentiate between various ideas.

**Evaluate (K5)** - Longer essay/Evaluation essay, Critique or justify with pros and cons.

Create (K6) – Check knowledge in specific or offbeat situations. Discussion.

#### **Mapping with Programme Outcomes:**

	PO 1	PO 2	PO 3	PO 4	PO 5	PO 6	PO 7	PO 8	PO 9	PO 10
CO 1	S	S	S	M	S	M	S	S	S	S
CO 2	S	S	S	L	M	M	S	S	S	S
CO 3	S	M	M	M	S	M	S	S	S	M
CO 4	S	M	M	L	S	L	S	S	S	M
CO 5	S	S	S	L	S	M	S	S	S	S

Subject Code: 23P4B18PW	PW: PROJECT WORK	Credits: 7
Semester: IV	WITH VIVA - VOCE	Maximum Marks: 100

Each candidate shall be required to take up a Project Work and submit it at the end of the final year. The Head of the Department shall assign the Guide who, in turn, will suggest the Project Work to the student in the beginning of the final year/semester. A copy of the Project Report will be submitted to the University/College through the Head of the Department on or before the date fixed by the University/College.

The Project will be evaluated by an internal and an external examiner nominated by the University/College. The candidate concerned will have to defend his/her Project through a Viva-voce.

## ASSESSMENT / EVALUATION / VIVA-VOCE

#### 1. PROJECT REPORT EVALUATION (Both Internal & External):

I. Plan of the Project

- 10 marks

- 50 marks

- 20 marks

II. Execution of the Plan / Collection of Data / Organisation of Materials / Fabrication of the Instruments / Experimental / Study / Hypothesis, Testing etc and

Presentation of the Report

III. Individual Initiative - 20 marks

2. VIVA-VOCE / INTERNAL & EXTERNAL

TOTAL -100 marks

## **PASSING MINIMUM**

Project Work	Viva-Voce 20 Marks	Dissertation 80 Marks
with Viva –	50% out of 20 Marks (i.e. 10 Marks)	50% out of 80 marks (i.e. 40 marks)
Voce		

A candidate shall be declared to have passed in the Project work if he/she gets not less than 50% in each of the Project Report and Viva voce and 50% in the aggregate of both the marks for Project Report and Viva-voce.

\*\*\*\*